GenCore version 5.1.6 Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on:

December 13, 2003, 04:37:24 ; Search time 1703.81 Seconds

(without alignments)

7460.491 Million cell updates/sec

US-09-852-261-5

Perfect score: 523

Sequence:

1 ggaccggagacgctctgcgg.....aaatacacaagtaaacattc 523

Scoring table:

IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched:

22781392 segs, 12152238056 residues

Total number of hits satisfying chosen parameters:

45562784

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

EST:*

- 1: em_estba:*
- 2: em_esthum:*
- 3: em estin:*
- 4: em estmu:*
- 5: em estov:*
- 6: em estpl:*
- 7: em estro:*
- 8: em_htc:*
- 9: gb est1:*
- 10: gb est2:*
- 11: gb htc:*
- 12: gb_est3:*
- 13: gb_est4:*
- 14: gb_est5:*
- 15: em estfun:*
- 16: em_estom:*
- 17: em gss_hum:*
- 18: em_gss_inv:*
- 19: em_gss_pln:*
- 20: em gss vrt:*
- 21: em_gss_fun:*
- 22: em gss mam: *
- 23: em_gss_mus:*
- 24: em gss pro:*
- 25: em gss rod:*
- 26: em gss phg:*
- 27: em gss vrl:*

28: gb_gss1:*
29: gb_gss2:*

용

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result			Query				
1	No.	Score	Match	Length	DB	ID	Description
	1	364.8	69.8	558	9	AI503976	AI503976 vm43d08.x
C	2	363	69.4	623	9	AW146128	AW146128 um37e10.x
C	3	348.2	66.6	549	9	AI169253	AI169253 EST215088
C	4	347	66.3	558	9	AI265629	AI265629 uj04b07.x
_	5	339.6	64.9	614	14	CD373004	CD373004 UI-R-GR0-
	6	339.2	64.9	816	9	AI119218	AI119218 ue94h02.y
	7	334.8	64.0	594	10	BF383724	BF383724 602044632
	8	334.4	63.9	796	14	CB959991	CB959991 AGENCOURT
С	9	322.2	61.6	499	9	AW495481	AW495481 UI-M-BH3-
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ACCESSION
            AI503976.1 GI:4401827
VERSION
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SOURCE
            Mus musculus (house mouse)
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            Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
               (bases 1 to 558)
REFERENCE
            Marra, M., Hillier, L., Kucaba, T., Martin, J., Beck, C., Wylie, T.,
  AUTHORS
            Underwood, K., Steptoe, M., Theising, B., Allen, M., Bowers, Y., Person
            ,B., Swaller,T., Gibbons,M., Pape,D., Harvey,N., Schurk,R., Ritter
            ,E., Kohn,S., Shin,T., Jackson,Y., Cardenas,M., McCann,R.,
            Waterston, R. and Wilson, R.
            The WashU-NCI Mouse EST Project 1999
  TITLE
  JOURNAL
            Unpublished
            Contact: Marra M/WashU-NCI Mouse EST Project 1999
COMMENT
            Washington University School of Medicine
            4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
            Tel: 314 286 1800
            Fax: 314 286 1810
            Email: mouseest@watson.wustl.edu
            This clone is available royalty-free through LLNL; contact the
            IMAGE Consortium (info@image.llnl.gov) for further information.
            MGI:565223
            This clone was previously sequenced on the 5' end only, this new
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VERSION
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REFERENCE
            (bases 1 to 623)
         Marra, M., Hillier, L., Kucaba, T., Martin, J., Beck, C., Wylie, T.,
 AUTHORS
         Underwood, K., Steptoe, M., Theising, B., Allen, M., Bowers, Y., Person
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,B., Swaller,T., Gibbons,M., Pape,D., Harvey,N., Schurk,R., Ritter
           ,E., Kohn,S., Shin,T., Jackson,Y., Cardenas,M., McCann,R.,
           Waterston, R. and Wilson, R.
 TITLE
           The WashU-NCI Mouse EST Project 1999
 JOURNAL
           Unpublished
COMMENT
           Contact: Marra M/WashU-NCI Mouse EST Project 1999
           Washington University School of Medicine
           4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
           Tel: 314 286 1800
           Fax: 314 286 1810
           Email: mouseest@watson.wustl.edu
           This clone is available royalty-free through LLNL; contact the
           IMAGE Consortium (info@image.llnl.gov) for further information.
           MGI:1006958
           Seg primer: custom primer used
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                   Location/Qualifiers
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                   double-stranded cDNA was ligated to a DraIII adaptor
                   [TGTTGGCCTACTGG], digested and cloned into distinct DraIII
                   sites of the pME18S-FL3 vector (5' site CACTGTGTG, 3' site
                   CACCATGTG). XhoI should be used to isolate the cDNA
                   insert. Size selection was performed to exclude fragments
                   <1.5kb. Library constructed by Dr. Sumio Sugano
                   (University of Tokyo Institute of Medical Science).
                   Custom primers for sequencing: 5' end primer
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TITLE	<pre>Kerlavage,A.R. and Adams,M.D. Rat Genome Project: Generation of a Rat EST (REST) Catalog & Rat</pre>
JOURNAL COMMENT	Gene Index Unpublished On Oct 6, 1998 this sequence version replaced gi:3705561. Other_ESTs: TC50779 Contact: Lee, NH The Institute for Genomic Research 9712, Medical Center Drive, Rockville, MD 20850, USA Tel: (301)-838-3529 Fax: (301)-838-0208 Email: nhlee@tigr.org Seq primer: M13-21.
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REFERENCE
            Marra, M., Hillier, L., Allen, M., Bowles, M., Dietrich, N., Dubuque, T.,
 AUTHORS
            Geisel, S., Kucaba, T., Lacy, M., Le, M., Martin, J., Morris, M.,
            Schellenberg, K., Steptoe, M., Tan, F., Underwood, K., Moore, B.,
            Theising, B., Wylie, T., Lennon, G., Soares, B., Wilson, R. and
            Waterston, R.
            The WashU-HHMI Mouse EST Project
 TITLE
            Unpublished
  JOURNAL
COMMENT
            Contact: Marra M/Mouse EST Project
            WashU-HHMI Mouse EST Project
            Washington University School of MedicineP
            4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
            Tel: 314 286 1800
            Fax: 314 286 1810
            Email: mouseest@watson.wustl.edu
            This clone is available royalty-free through LLNL; contact the
            IMAGE Consortium (info@image.llnl.gov) for further information.
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                     ligated to a DraIII adaptor [TGTTGGCCTACTGG], digested
                     and cloned into distinct DraIII sites of the pME18S-FL3
                     vector (5' site CACTGTGTG, 3' site CACCATGTG). XhoI should
                     be used to isolate the cDNA insert. Size selection was
                     performed to exclude fragments <1.5kb. Library
                     constructed by Dr. Sumio Sugano (University of Tokyo
                     Institute of Medical Science). Custom primers for
                     sequencing: 5' end primer CTTCTGCTCTAAAAGCTGCG and 3' end
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AI265629.1 GI:3883787

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VERSION
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         Rattus.
REFERENCE
           (bases 1 to 614)
         Bonaldo, M.F., Lennon, G. and Soares, M.B.
 AUTHORS
 TITLE
         Normalization and subtraction: two approaches to facilitate gene
         discovery
 JOURNAL
         Genome Res. 6 (9), 791-806 (1996)
 MEDLINE
         97044477
  PUBMED
         8889548
```

```
Contact: Soares, MB
COMMENT
           Coordinated Laboratory for Computational Genomics
           University of Iowa
           375 Newton Road , 4156 MEBRF, Iowa City, IA 52242, USA
           Tel: 319 335 8250
           Fax: 319 335 9565
           Email: bento-soares@uiowa.edu
           Tissue Procurement: James Lin, University of Iowa
            cDNA Library preparation: Dr. M. Bento Soares, University of Iowa
            cDNA Library Arrayed by: Dr. M. Bento Soares, University of Iowa
            DNA Sequencing by: Dr. M. Bento Soares, University of Iowa
            Clone Distribution: Distribution information can be found at
           http://genome.uiowa.edu/distribution/rat.html
           Seq primer: M13 REVERSE.
                    Location/Qualifiers
FEATURES
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                    /strain="Sprague-Dawley"
                    /db xref="taxon:10116"
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                    /tissue type="Whole embryo"
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                    /lab host="DH10B (Life Technologies) (T1 phage resistant)"
                    /clone lib="UI-R-GR0"
                    /note="Vector: pYX-Asc; Site 1: EcoR I; Site 2: Not I;
                    UI-R-GRO is a cDNA library containing the following
                    tissue(s): rat whole embryo 13dpc. The library was
                    constructed according to Bonaldo, Lennon and Soares,
                    Genome Research, 6:791-806, 1996. Denatured RNA was size
                    fractionated on a 1% agarose gel. First strand cDNA
                    synthesis was primed with oligo-dT primer containing a Not
                    I site. Double strand cDNA was size selected according to
                    mRNA size fraction, ligated with EcoR I adaptor, digested
                    with NotI and then cloned directionally into pYX-Asc
                    vector. The library tag sequence located between the Not I
                    site and the polyA tail is CATCTCTACT. This library was
                    created for the University of Iowa Program for Rat Gene
                    Discovery and Mapping (Val Sheffield, Bento Soares and Tom
                    Casavant)."
                        168 c
BASE COUNT
                                         119 t
                                                    2 others
               171 a
                                154 q
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  Query Match
                         64.9%;
                                                     Length 614;
                         81.4%; Pred. No. 1.9e-74;
  Best Local Similarity
  Matches 393; Conservative
                               0; Mismatches
                                                90;
                                                   Indels
                                                              0; Gaps
                                                                          0;
           1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGGAGAC 60
Qу
              Db
         116 GGACCAGAGACCCTTTGCGGGGCTGAGCTGGTGGACGCTCTTCAGTTCGTGTGTGGACCA 175
          61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGCCACCTCAG 120
Qу
              176 AGGGGCTTTTACTTCAACAAGCCCACAGGCTATGGCTCCAGCATTCGGAGGGCACCACAG 235
Db
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Qу
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MGI:936407 Seq primer: custom primer used High quality sequence stop: 473. Location/Qualifiers **FEATURES** 1. .816 source /organism="Mus musculus" /mol type="mRNA" /strain="C57BL" /db xref="taxon:10090" /clone="IMAGE:1498803" /dev stage="embryo, 14 dpc" /lab host="DH10B" /clone lib≈"Sugano mouse embryo mewa" /note="Vector: pME18S-FL3; Site_1: DraIII (CACTGTGTG); Site 2: DraIII (CACCATGTG); 1st strand cDNA was primed with an oligo(dT) primer [ATGTGGCCTTTTTTTTTTTTT]; double-stranded cDNA was ligated to a DraIII adaptor [TGTTGGCCTACTGG], digested and cloned into distinct DraIII sites of the pME18S-FL3 vector (5' site CACTGTGTG, 3' site CACCATGTG). XhoI should be used to isolate the cDNA insert. Size selection was performed to exclude fragments <1.5kb. Library constructed by Dr. Sumio Sugano (University of Tokyo Institute of Medical Science). Custom primers for sequencing: 5' end primer CTTCTGCTCTAAAAGCTGCG and 3' end primer CGACCTGCAGCTCGAGCACA." BASE COUNT 172 g 8 others 230 a 219 c 187 t ORIGIN 64.9%; Score 339.2; DB 9; Query Match Length 816; 81.2%; Pred. No. 2.5e-74; Best Local Similarity Matches 389; Conservative 0; Mismatches 90; Indels 0;

Qу	1	GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 60
Db	323	GGACCAGAGACCCTTTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGACCG 382
Qy	61	AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
Db	383	AGGGGCTTTTACTTCAACAAGCCCACAGGCTATGGCTCCAGCATTCGGAGGGCACCTCAG 442
Qy	121	ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Db	443	ACAGGCATTGTGGATGAGTGTTGCTTCCGGAGCTGTGATCTGAGGAGACTGGAGATGTAC 502
Qу	181	TGTGCACCCCTCAAGCCGGCAAAGGCAGCCCGCTCCGTCCG
Db	503	TGTGCCCCACTGAAGCCTACAAAAGCAGCCCGCTCTATCCGTGCCCAGCGCCACACTGAC 562
Qу	241	ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
Db	563	ATGCCCAAGACTCAGAAGTCCCCGTCCCTATCGACAAACAA
Qy	301	AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
Db	623	AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGAAGTGCANGAAACAAGACCTA 682

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Qу
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             683 CAGAATGTANGAGGAGCCTNCCACGGAGCAGAANATGCCACATCACCGCANGATCCTTTG 742
Db
         421 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATT 479
Qу
                               743 CTGCTTGAGCAACCTGCANAACATCGAAACACCTACCAAATAACATNTATAAGTCCAAT 801
Db
RESULT 7
BF383724
                                   594 bp
                                             mRNA
                                                     linear
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LOCUS
           BF383724
           602044632F1 NCI CGAP Li9 Mus musculus cDNA clone IMAGE:4194295 5',
DEFINITION
           mRNA sequence.
ACCESSION
           BF383724
VERSION
           BF383724.1 GI:11365029
KEYWORDS
           EST
SOURCE
           Mus musculus (house mouse)
  ORGANISM Mus musculus
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE
              (bases 1 to 594)
           NIH-MGC http://mgc.nci.nih.gov/.
 AUTHORS
           National Institutes of Health, Mammalian Gene Collection (MGC)
  TITLE
  JOURNAL
           Unpublished
COMMENT
           Contact: Robert Strausberg, Ph.D.
           Email: cgapbs-r@mail.nih.gov
           Tissue Procurement: Jeffrey E. Green, M.D.
            cDNA Library Preparation: Life Technologies, Inc.
            cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
            DNA Sequencing by: Incyte Genomics, Inc.
            Clone distribution: MGC clone distribution information can be
           found through the I.M.A.G.E. Consortium/LLNL at:
           http://image.llnl.gov
           Plate: LLAM9527 row: p column: 08
           High quality sequence stop: 589.
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                    /clone="IMAGE:4194295"
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                    /note="Organ: liver; Vector: pCMV-SPORT6; Site 1: NotI;
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                    Technologies. Note: this is a NCI CGAP Library."
BASE COUNT
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                        162 c
                                142 g
                                         115 t
ORIGIN
                         64.0%; Score 334.8; DB 10; Length 594;
  Query Match
  Best Local Similarity 82.0%; Pred. No. 2.9e-73;
  Matches 400; Conservative
                               0; Mismatches
                                                     Indels
                                                82;
                                                                          1;
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Qу

Total	Db									
Oy 16 CCGCCACAGAGACCCCCTACCACAAACAAACAACAACAACAACACCCCCTAGAGAGAG	Qy	76 AACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAGACAGGCATCGTGGAT 135								
	Db									
Db 227 GAGTGTTCCTTCCGGAGCTGTGATCTGAGGAGATGTACTGTGCCCCACTGAAG 286 Qy 196 CCGGCAAAGGCAGCCCGCTCCGTCCGTGCCAGCCCACACCGACATGCCCAAGACTCAG 255	Qy									
	Db									
Qy 256 AAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGGAGAAAGGAAAGT 315	QУ	196 CCGGCAAAGGCAGCCCGCTCCGTCCGTGCCCAGCGCCACACCGACATGCCCAAGACTCAG 255								
Db 347 AAGTCCCCGTCCCTATCGACAAACAAGAAACCGAAGCTGCAAAGGAAAGGAAAGGAAGT 406 Qy 316 ACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTACAGGATGTAGGAAGA 375	Db	287 CCTACAAAAGCAGCCCGCTCTATCCGTGCCCAGCGCCACACTGACATGCCCAAGACTCAG 346								
Db 347 AAGTCCCCGTCCCTATCGACAAACAAGAAACGAAGCTGCAAAGGAAGG	Qy									
Db 407 ACATTTGAAGAACACAAGTAGAGGAAGTGCAGGAAACAAGACCTACAGAATGTAGGAGGA 466 Qy 376 CCCTTCTGAGGAGTGAAGAAGAGGCCACCGCAGGACCCTTTGCTCTGCACAGTTACC 435	Db									
Qy 376 CCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTGCTCTGCACAGTTACC 435	Qy	316 ACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTACAGGATGTAGGAAGA 375								
Db 467 GCCTCCCACGGAGCAGAAAATGCCACATCACCGCAGGATCCTTTGCTGCTTGAGCAACCT 526 Qy 436 TGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTTCAAAGATGG 489	Db	407 ACATTTGAAGAACACAAGTAGAGGAAGTGCAGGAAACAAGACCTACAGAATGTAGGAGGA 466								
Qy 436 TGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTTCAAAGATGG 489	Qy									
Db 527 GCAAAACATCGAAACACCTACCAAATAACAATAATAAGTCCAATAACATTACAAAGATGG 586 Qy 490 CATTTCCC 497	Db	467 GCCTCCCACGGAGCAGAAAATGCCACATCACCGCAGGATCCTTTGCTGCTTGAGCAACCT 526								
Qy 490 CATTTCCC 497 Db 587 GCATTGCC 594 RESULT 8 CB959991 LOCUS CB959991 796 bp mRNA linear EST 29-APR-2003 DEFINITION AGENCOURT 13888044 NIH MGC 147 Homo sapiens cDNA clone	QУ	436 TGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTTCAAAGATGG 489								
RESULT 8 CB959991 LOCUS CB959991 796 bp mRNA linear EST 29-APR-2003 DEFINITION AGENCOURT_13888044 NIH_MGC_147 Homo sapiens cDNA clone	Db	527 GCAAAACATCGAAACACCTACCAAATAACAATAAGTCCAATAACATTACAAAGATGG 586								
RESULT 8 CB959991 LOCUS CB959991 796 bp mRNA linear EST 29-APR-2003 DEFINITION AGENCOURT_13888044 NIH_MGC_147 Homo sapiens cDNA clone	QУ	490 CATTTCCC 497								
CB959991 LOCUS CB959991 796 bp mRNA linear EST 29-APR-2003 DEFINITION AGENCOURT_13888044 NIH_MGC_147 Homo sapiens cDNA clone IMAGE:30341081 5', mRNA sequence. ACCESSION CB959991 VERSION CB959991.1 GI:30216107 KEYWORDS EST. SOURCE Homo sapiens (human) ORGANISM Homo sapiens Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;	Db	587 GCATTGCC 594								
KEYWORDS EST. SOURCE Homo sapiens (human) ORGANISM Homo sapiens Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;	CB959991 LOCUS DEFINITIO	ON AGENCOURT_13888044 NIH_MGC_147 Homo sapiens cDNA clone IMAGE:30341081 5', mRNA sequence. CB959991								
ORGANISM Homo sapiens Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;		EST.								
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;		Homo sapiens								
		Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.								
REFERENCE 1 (bases 1 to 796) AUTHORS NIH-MGC http://mgc.nci.nih.gov/.	AUTHORS	NIH-MGC http://mgc.nci.nih.gov/.								
TITLE National Institutes of Health, Mammalian Gene Collection (MGC) JOURNAL Unpublished	JOURNAI	_ Unpublished								
COMMENT Contact: Robert Strausberg, Ph.D. Email: cgapbs-r@mail.nih.gov	COMMENT	Email: cgapbs-r@mail.nih.gov								
Tissue Procurement: Dr. Stefan Hansson cDNA Library Preparation: Michael J. Brownstein (NHGRI) with help		Tissue Procurement: Dr. Stefan Hansson cDNA Library Preparation: Michael J. Brownstein (NHGRI) with help and advice from Piero Carninci (RIKEN)								
		and advice from Piero Carninci (RIKEN)								

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cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
          DNA Sequencing by: Agencourt Bioscience Corporation
          Clone distribution: MGC clone distribution information can be
          found through the I.M.A.G.E. Consortium/LLNL at:
          http://image.llnl.gov
          Plate: NDAM371 row: p column: 18
          High quality sequence stop: 707.
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                 /lab host="DH10B TonA"
                 /clone lib="NIH MGC 147"
                 /note="Organ: placenta; Vector: pBluescriptR; Site 1:
                 all-XhoI; Site 2: BamH; Oligo-dT primed using primer
                 5'-TTTTTTTTTTTTTTVN-3', size-selected for average
                 insert size 2.3 kb and normalized to ROT 5. This is a
                 primary library enriched for full-length clones and
                 constructed using the Cap-trapper method (Carninci, in
                 preparation). Library constructed by M. Brownstein
                 (NIMH/NHGRI, National Institutes of Health). Note: This is
                 a NIH MGC library."
                     197 c
BASE COUNT
             224 a
                            191 q
                                    184 t
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                     84.6%; Pred. No. 3.9e-73;
 Best Local Similarity
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            Db
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Qу
            Db
        240 AGGGGCTTTTATTTCAACAAGCCCACAGGGTATGGCTCCAGCAGTCGGAGGGCGCCTCAG 299
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            Db
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            420 ATGCCCAAGACCCAG-
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                  Db
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Qу

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         479 TTCAAAGAT-GGCATTTCCCCCAATGAAATACACAAGTAAACATTC 523
Qу
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RESULT 9
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                                                            EST 24-FEB-2000
LOCUS
           AW495481
           UI-M-BH3-auy-g-11-0-UI.s1 NIH BMAP M S4 Mus musculus cDNA clone
DEFINITION
           UI-M-BH3-auy-g-11-0-UI 3', mRNA sequence.
ACCESSION
           AW495481
           AW495481.1 GI:7065762
VERSION
           EST.
KEYWORDS
           Mus musculus (house mouse)
SOURCE
 ORGANISM Mus musculus
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE
              (bases 1 to 499)
 AUTHORS
           Bonaldo, M.F., Lennon, G. and Soares, M.B.
           Normalization and subtraction: two approaches to facilitate gene
 TITLE
           Genome Res. 6 (9), 791-806 (1996)
 JOURNAL
 MEDLINE
           97044477
  PUBMED
           8889548
COMMENT
           Contact: Chin, H
           National Institute of Mental Health
           6001 Executive Blvd. Room 7N-7190, MSC 9643, Bethesda, MD
           20892-9643, USA
           Tel: 301 443 1706
           Fax: 301 443 9890
           Email: mEST@mail.nih.gov
           The sequence contained an oligo-dT track that was present in the
           oligonucleotide that was used to prime the synthesis of first
           strand cDNA and therefore this may represent a bonafide poly A
           tail. The sequence tag present in the cDNA between the NotI site
           and the oligo-dT track served to identify it as a clone from the
           normalized pineal glands library cDNA Library Preparation: M.B.
           Soares Lab Clone distribution: Researchers may obtain BMAP cDNA
           clones from RESEARCH GENETICS. It should be noted that Bento Soares
           is generating a small number of additional specialized
           non-redundant arrays of BMAP cDNAs whose availability will be
           considered under appropriate and limited collaborative arrangements
           Seg primer: M13 Forward
           POLYA=Yes.
                   Location/Qualifiers
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/db xref="taxon:10090"

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                 /lab host="DH10B (Life Technologies)"
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                 /note="Vector: pT7T3D-Pac (Pharmacia) with a modified
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                 NIH BMAP M S4 library is a subtracted library of a series,
                 ultimately derived from a mixture of individually tagged
                 normalized libraries from ten regions of the mouse brain
                 (cerebellum, brain stems, olfactory bulbs, hypothalamus,
                 cortex, amygdala, basal ganglia, pineal gland, striatum,
                 hipoccampus) after a series of subtractions to reduce the
                 representation of cDNAs from which ESTs had already been
                 generated. The following serially subtracted libraries
                 were generated in this process: NIH BMAP M S4,
                 NIH BMAP M S3.3, NIH BMAP M S3.2, NIH BMAP M S3.1,
                 NIH BMAP M S2, NIH BMAP M S1. The subtracted library
                 (NIH_BMAP_M_S4) was constructed as follows: PCRamplified
                 cDNA inserts from NIH BMAP M S3.3, NIH BMAP M S3.2, and
                 NIH BMAP M S3.1 clones from which 3' ESTs had been derived
                 was used as a driver in a hybridization with a pool of
                 the NIH BMAP M S3.3, NIH BMAP M S3.2, and NIH BMAP M S3.1
                 libraries in the form of single-stranded circles. The
                 remaining single-stranded circles (subtracted library)
                 was purified by hydroxyapatite column chromatography,
                 converted to double-stranded circles and electroporated
                 into DH10B bacteria (LifeTechnologies) to generate the
                 NIH BMAP M S4 library. This procedure has been previously
                 described (Bonaldo, Lennon and Soares, Genome Research
                 6:791-806, 1996)
                 TAG LIB=NIH BMAP M S4
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                 TAG SEQ=CAGAC"
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                            124 g
                                    177 t
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Best Local Similarity 82.3%; Pred. No. 4e-70;
Matches 396; Conservative
                           0; Mismatches
                                                                   2;
                                           78; Indels
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                    499 TGTGTGGACCGAGGGGTTTTACTTCAACAAGCCCACAGGCTATGGCTCCAGCATTCGGA 440
       110 GGGCACCTCAGACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGC 169
           439 GGGCACCTCAGACAGGCATTGTGGATGAGTGTTGCTTCCGGAGCTGTGATCTGAGGAGAC 380
       170 TGGAGATGTACTGTGCACCCCTCAAGCCGGCAAAGGCAGCCCGCTCCGTCCCTGCCCAGC 229
           379 TGGAGATGTACTGTGCCCCACTGAAGCCTACAAAAGCAGCCCGCTCTATCCGTGCCCAGC 320
       230 GCCACACCGACATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGA 289
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BASE COUNT

ORIGIN

QУ

Db

Qу

Db

Qу

Db

Qу

Db

Qу

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QУ
             Db
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         410 AGGACCCTTTGCTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCA-----AAAAA 463
QУ
             139 AGGATCCTTTGCTGCTTGAGCAACCTGCAAAACATCGAAACACCTACCAAATAACAATAA 80
Db
         464 TAAGTTTGATCACATTTCAAAGAT-GGCATTTCCCCCCAATGAAATACACAAGTAAACATT 522
QУ
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Db
         523 C 523
Qу
Db
          19 C 19
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AI876493/c
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           AI876493
                                  642 bp
                                           mRNA
                                                   linear
                                                           EST 21-JUL-1999
           uj59b10.x1 Sugano mouse liver mlia Mus musculus cDNA clone
DEFINITION
           IMAGE:1924219 3' similar to gb:X57025 rna1 INSULIN-LIKE GROWTH
           FACTOR IA PRECURSOR (HUMAN); gb:X04482 Mouse mRNA for
           preproinsulin-like growth factor IB (MOUSE);, mRNA sequence.
ACCESSION
           AI876493
VERSION
           AI876493.1 GI:5550542
KEYWORDS.
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SOURCE
           Mus musculus (house mouse)
  ORGANISM
           Mus musculus
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE
              (bases 1 to 642)
           Marra, M., Hillier, L., Kucaba, T., Martin, J., Beck, C., Wylie, T.,
 AUTHORS
           Underwood, K., Steptoe, M., Theising, B., Allen, M., Bowers, Y., Person
           ,B., Swaller,T., Gibbons,M., Pape,D., Harvey,N., Schurk,R., Ritter
           ,E., Kohn,S., Shin,T., Jackson,Y., Cardenas,M., McCann,R.,
           Waterston, R. and Wilson, R.
  TITLE
           The WashU-NCI Mouse EST Project 1999
  JOURNAL
           Unpublished
COMMENT
           Contact: Marra M/WashU-NCI Mouse EST Project 1999
           Washington University School of Medicine
           4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
           Tel: 314 286 1800
           Fax: 314 286 1810
           Email: mouseest@watson.wustl.edu
           This clone is available royalty-free through LLNL; contact the
           IMAGE Consortium (info@image.llnl.gov) for further information.
           MGI:980511
           Seq primer: custom primer used
           High quality sequence stop: 257.
FEATURES
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                 was primed with an oligo(dT) primer
                 [ATGTGGCCTTTTTTTTTTTTTT]; double-stranded cDNA was
                 ligated to a DraIII adaptor [TGTTGGCCTACTGG], digested
                 and cloned into distinct DraIII sites of the pME18S-FL3
                 vector (5' site CACTGTGTG, 3' site CACCATGTG). XhoI should
                 be used to isolate the cDNA insert. Size selection was
                 performed to exclude fragments <1.5kb. Library
                 constructed by Dr. Sumio Sugano (University of Tokyo
                 Institute of Medical Science). Custom primers for
                 sequencing: 5' end primer CTTCTGCTCTAAAAGCTGCG and 3' end
                 primer CGACCTGCAGCTCGAGCACA."
BASE COUNT
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                           175 q
ORIGIN
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                     61.3%;
                                            Length 642;
 Best Local Similarity
                     80.1%; Pred. No. 9.6e-70;
 Matches 403; Conservative
                          0; Mismatches
                                            Indels
                                                       Gaps
                                                              2;
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Qу
           Db
        503 GACCAGAGACCCTTTGCGGGGCTGAGCTGGTGGATGCTCTTCAGGTCGTGTGTGGACCGA 444
        62 GGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAGA 121
Qу
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        443 GGGGCTTTTCTTCAACAAGGCCACAGGCTATGGCTCCAGCATTTGGAGGGCACCTCAGA 384
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           Db
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Qу
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              Db
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                                                       linear
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            UI-CF-EC1-abj-k-24-0-UI.sl UI-CF-EC1 Homo sapiens cDNA clone
DEFINITION
            UI-CF-EC1-abj-k-24-0-UI 3', mRNA sequence.
            BM984670
ACCESSION
VERSION
            BM984670.1 GI:19610417
KEYWORDS
            EST.
            Homo sapiens (human)
SOURCE
 ORGANISM
           Homo sapiens
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            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
               (bases 1 to 673)
REFERENCE
            Bonaldo, M.F., Lennon, G. and Soares, M.B.
 AUTHORS
            Normalization and subtraction: two approaches to facilitate gene
  TITLE
            discovery
            Genome Res. 6 (9), 791-806 (1996)
  JOURNAL
  MEDLINE
            97044477
   PUBMED
            8889548
COMMENT
            Contact: McCray, PB
            McCray Lab
            University of Iowa
            2024 University of Iowa Med Labs, Iowa City, IA 52242, USA
            Tel: 319 356 4866
            Fax: 319 356 7171
            Email: paul-mccray@uiowa.edu
            Tissue Procurement: Dr. M. J. Welsh, University of Iowa
             cDNA Library preparation: Dr. M. Bento Soares, University of Iowa
             cDNA Library Arrayed by: Dr. M. Bento Soares, University of Iowa
            DNA Sequencing by: Dr. M. Bento Soares, University of Iowa
             Clone Distribution: Researchers may obtain clones from Research
            Genetics (www.resgen.com) or from Open Biosystems
            (www.openbiosystems.com).
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            POLYA=Yes.
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                     /clone lib="UI-CF-EC1"
                     /note="Organ: Lung; Vector: pT7T3-Pac (Pharmacia) with a
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                     UI-CF-EC1 is a normalized cDNA library containing the
                     following tissue(s): Normal lung from adult and from fetal
```

day 64, day 87, week 19 and week 42. The library was

constructed according to Bonaldo, Lennon and Soares, Genome Research, 6:791-806, 1996. First strand cDNA synthesis was primed with an oligo-dT primer containing a Not I site. Double stranded cDNA was ligated to an EcoR I adaptor, digested with Not I, and cloned directionally into pT7T3-Pac vector. The oligonucleotide used to prime the synthesis of first-strand cDNA contains a library tag sequence that is located between the Not I site and the (dT)18 tail. The sequence tag for this library is AAGTGCTTAC.

TAG LIB=UI-CF-EC1

TAG_TISSUE=Normal Lung Epithelial Cells Tissue nos 369-371 and 380-383

TAG SEQ=AAGTGCTTAC"

BASE COUNT ORIGIN 152 a 164 c 169 g 188 t

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                                                     linear
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DEFINITION
           clone IMAGE:1849953 3' similar to gb:X57025 rna1 INSULIN-LIKE
           GROWTH FACTOR IA PRECURSOR (HUMAN);, mRNA sequence.
ACCESSION
           AI248089
           AI248089.1 GI:3843486
VERSION
KEYWORDS
           EST.
SOURCE
           Homo sapiens (human)
 ORGANISM
           Homo sapiens
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           Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
           1 (bases 1 to 575)
REFERENCE
           NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
 AUTHORS
           National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
 TITLE
           Tumor Gene Index
           Unpublished
  JOURNAL
           Contact: Robert Strausberg, Ph.D.
COMMENT
           Email: cgapbs-r@mail.nih.gov
           This clone is available royalty-free through LLNL; contact the
           IMAGE Consortium (info@image.llnl.gov) for further information.
                               Std Error: 0.00
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                    This is a subtracted version of the original Soares fetal
                    liver spleen 1NFLS library.
                                                 1st strand cDNA was primed
                    with a Pac I - oligo(dT) primer [5'
                    double-stranded cDNA was ligated to Eco RI adaptors
                    (Pharmacia), digested with Pac I and cloned into the Pac I
                    and Eco RI sites of the modified pT7T3 vector. Library
                    went through one round of normalization. Library
                    constructed by Bento Soares and M. Fatima Bonaldo."
               135 a
                        152 c
                                 131 g
                                         156 t
                                                    1 others
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  Best Local Similarity
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Qу
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551 TGCGGGGCTGAGCTGGTGNATGCTCTTCAGTTCGTGTGAAGACAGGGGCTTTTATTTC 492

Db

Qy	76 AACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAGACAGGCATCGTGGAT 135
Db	491 AACAAGCCCACAGGGTATGGCTCCAGCAGTCGGAGGGCGCCTCAGACAGGCATCGTGGAT 432
QУ	136 GAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTACTGTGCACCCCTCAAG 195
Db	431 GAGTGCTTCCGGAGCTGTGATCTAAGGAGGCTGGAGATGTATTGCGCACCCCTCAAG 372
Qy	196 CCGGCAAAGGCAGCCCGCTCCGTCCCTGCCCAGCGCCACACCGACATGCCCAAGACTCAG 255
Db a	371 CCTGCCAAGTCAGCTCGCTCTGTCCGTGCCCAGCGCCACACCGACATGCCCAAGACCCAG 312
Qу	256 AAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGGAGAAGGAAAGGAAGT 315
Db	 311AAGGAAGT 304
Qу	316 ACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTACAGGATGTAGGAAGA 375
Db	
Qу	376 CCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTGCTCTGCAC-AGTTAC 434
Db	
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Db	
Qy	493 TTCCCCCAATGAAATACACAAGTAAACATTC 523
Db	
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	The Institute for Genomic Research 9712, Medical Center Drive, Rockville, MD 20850, USA

```
Tel: (301)-838-3529
         Fax: (301) -838-0208
         Email: nhlee@tigr.org
         Seg primer: M13-21.
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QУ
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           408 GGGCATTGTGGATGAGTGTTGCTCCCGGAGCTGTGATCTGAGGAGGTTGGAGATGTACTG 349
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                       Dh
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RESULT 14 AA542914/c

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Db 221AAGGAAGTACATTTGAAGAACGCAAGTAGAGGGGAGTGCAGGAAACAAGAACT 170 Qy 360 ACAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGACGCACCGCAGGACCCTTT 419
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Db 169 ACAGGATGTAGGAAGACCCTCCTGAGGAGTGACATGCCACCGCAGGATCCTTT 110 Qy 420 GCTCTGCAC-AGTTACCTG-TAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACA 477
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Db 109 GCTCTGCACGAGTTACCTGTTAAACTTTGGAACACCTACCAAAAAATAAGTTTGATAACA 50 Qy 478 TTTCAAAGAT-GGCATTTCCCCCCAATGAAATACACAAGTAAACATTC 523
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Db 49 TTTAAAAGATGGGCGTTTCCCCCAATGAAATACACAAGTAAACATTC 3 RESULT 15 A1604642 LOCUS A1604642 882 bp mRNA linear EST 21-APR-1999 DEFINITION vm43d08.yl Stratagene mouse diaphragm (#937303) Mus musculus cDNA
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VERSION A1604642.1 GI:4613809 KEYWORDS EST.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. REFERENCE 1 (bases 1 to 882)
AUTHORS Marra,M., Hillier,L., Kucaba,T., Martin,J., Beck,C., Wylie,T., Underwood,K., Steptoe,M., Theising,B., Allen,M., Bowers,Y., Person ,B., Swaller,T., Gibbons,M., Pape,D., Harvey,N., Schurk,R., Ritter ,E., Kohn,S., Shin,T., Jackson,Y., Cardenas,M., McCann,R., Waterston,R. and Wilson,R.
TITLE The WashU-NCI Mouse EST Project 1999
JOURNAL Unpublished COMMENT Contact: Marra M/WashU-NCI Mouse EST Project 1999 Washington University School of Medicine

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REFERENCE ·
            1
               (bases 1 to 498)
  AUTHORS
            NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
            National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
  TITLE
            Tumor Gene Index
            Unpublished
  JOURNAL
            Contact: Robert Strausberg, Ph.D.
COMMENT
            Email: cgapbs-r@mail.nih.gov
            Tissue Procurement: Michael J. Brownstein, M.D., Ph.D., Michael R.
            Emmert-Buck, M.D., Ph.D.
            cDNA Library Preparation: M. Bento Soares, Ph.D.
            cDNA Library Arrayed by: Greg Lennon, Ph.D.
            DNA Sequencing by: Washington University Genome Sequencing Center
            Clone distribution: NCI-CGAP clone distribution information can be
            found through the I.M.A.G.E. Consortium/LLNL at:
            www-bio.llnl.gov/bbrp/image/image.html
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                    a Not I - oligo(dT) primer. Double-stranded cDNA was
                     ligated to Eco RI adaptors (Pharmacia), digested with Not
                     I and cloned into the Not I and Eco RI sites of the
                    modified pT7T3 vector. Library is not normalized. Library
                    was constructed by Bento Soares and M. Fatima Bonaldo. "
BASE COUNT
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 Matches 440; Conservative
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Qу
                          Dh
          476 GGACCGGAGAACTTTTGCGGGGCTTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGA 417
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```
Tel: 314 286 1800
          Fax: 314 286 1810
          Email: mouseest@watson.wustl.edu
         This clone is available royalty-free through LLNL; contact the
          IMAGE Consortium (info@image.llnl.gov) for further information.
         MGI:565223
         This read is a RESEQUENCE of a previously sequenced mouse clone
         This read has been verified (found to hit its original self in the
         correct orientation)
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         High quality sequence stop: 361.
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BASE COUNT
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Qу
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                 Db
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Qу
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4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA

Dp 6.	7 GAGAGGGAAGGGAGTACATTTGAGGAACACANGTNGAGGAAGTGCANGAAACAAGACCTA 736
Qy 3	1 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Db 73	
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Search completed: December 13, 2003, 07:29:51

Job time : 1704.81 secs

GenCore version 5.1.6 Copyright (c) 1993 - 2003 Compugen Ltd.

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December 13, 2003, 05:41:20; Search time 2336.77 Seconds

(without alignments)

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US-09-852-261-5

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Minimum DB seq length: 0

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Maximum Match 100%

Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

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VERSION
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REFERENCE
  AUTHORS
           Goldspink, G.R. and Johnson, I.R.
  TITLE
           Use of the insulin-like-growth factor i isoform mgf for the
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REFERENCE
 AUTHORS
        Goldspink, G.D. and Terenghi, G.B.
 TITLE
        Repair of nerve damage
 JOURNAL
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REFERENCE
         Goldspink, G.R. and Johnson, I.R.
 AUTHORS
         Use of the insulin-like-growth factor i isoform mgf for the
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 AUTHORS
          Goldspink, G.D. and Terenghi, G.B.
          Repair of nerve damage
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REFERENCE
 AUTHORS
         Goldspink, G.R. and Johnson, I.R.
 TITLE
         Use of the insulin-like-growth factor i isoform mgf for the
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ACCESSION AX300791

VERSION AX300791.1 GI:17382072

KEYWORDS

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ORGANISM Oryctolagus cuniculus

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         Repair of nerve damage
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 JOURNAL
         University College London (GB); East Grinstead Medical Research
         Trust (GB)
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LOCUS
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DEFINITION
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ACCESSION
            BC012409
VERSION
            BC012409.1 GI:15214568
KEYWORDS
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SOURCE
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REFERENCE
               (bases 1 to 1536)
            Strausberg, R.L., Feingold, E.A., Grouse, L.H., Derge, J.G.,
  AUTHORS
            Klausner, R.D., Collins, F.S., Wagner, L., Shenmen, C.M., Schuler, G.D.,
            Altschul, S.F., Zeeberg, B., Buetow, K.H., Schaefer, C.F., Bhat, N.K.,
            Hopkins, R.F., Jordan, H., Moore, T., Max, S.I., Wang, J., Hsieh, F.,
            Diatchenko, L., Marusina, K., Farmer, A.A., Rubin, G.M., Hong, L.,
            Stapleton, M., Soares, M.B., Bonaldo, M.F., Casavant, T.L.,
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            Carninci, P., Prange, C., Raha, S.S., Loquellano, N.A., Peters, G.J.,
            Abramson, R.D., Mullahy, S.J., Bosak, S.A., McEwan, P.J.,
            McKernan, K.J., Malek, J.A., Gunaratne, P.H., Richards, S.,
            Worley, K.C., Hale, S., Garcia, A.M., Gay, L.J., Hulyk, S.W.,
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            Dickson, M.C., Rodriguez, A.C., Grimwood, J., Schmutz, J., Myers, R.M.,
            Butterfield, Y.S., Krzywinski, M.I., Skalska, U., Smailus, D.E.,
            Schnerch, A., Schein, J.E., Jones, S.J. and Marra, M.A.
  TITLE
            Generation and initial analysis of more than 15,000 full-length
            human and mouse cDNA sequences
  JOURNAL
            Proc. Natl. Acad. Sci. U.S.A. 99 (26), 16899-16903 (2002)
  MEDLINE
            22388257
   PUBMED
            12477932
               (bases 1 to 1536)
REFERENCE
  AUTHORS
            Strausberg, R.
  TITLE
            Direct Submission
  JOURNAL
            Submitted (15-AUG-2001) National Institutes of Health, Mammalian
            Gene Collection (MGC), Cancer Genomics Office, National Cancer
            Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590,
            USA
  REMARK
            NIH-MGC Project URL: http://mgc.nci.nih.gov
COMMENT
            Contact: MGC help desk
            Email: cgapbs-r@mail.nih.gov
            Tissue Procurement: Jeffrey E. Green, M.D.
            cDNA Library Preparation: Life Technologies, Inc.
            cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
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DNA Sequencing by: Baylor College of Medicine Human Genome

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Center code: BCM-HGSC
          Web site: http://www.hgsc.bcm.tmc.edu/cdna/
          Contact: amg@bcm.tmc.edu
          Gunaratne, P.H., Garcia, A.M., Lu, X., Hulyk, S.W., Loulseged, H.,
          Kowis, C.R., Sneed, A.J., Martin, R.G., Muzny, D.M., Nanavati,
          A.N., Gibbs, R.A.
          Clone distribution: MGC clone distribution information can be found
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Sequencing Center

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DEFINITION Rat mRNA (clone IGF1AB2) for insulin-like growth factor I.
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ACCESSION
          X06108.1 GI:56426
VERSION
KEYWORDS
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SOURCE
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REFERENCE
          1 .
 AUTHORS
          Shimatsu, A. and Rotwein, P.
 TITLE
          Sequence of Two Rat Insulin-like Growth Factor I mRNAs Differing
          Within the 5' Untranslated Region
 JOURNAL
          Nucleic Acids Res. 15 (1987) In press
REFERENCE
            (bases 1 to 798)
 AUTHORS
          Rotwein, P.
 TITLE
          Direct Submission
 JOURNAL
          Submitted (21-OCT-1987) Rotwein P., Washington University, School
          of Medicine, 660 South Euclid Avenue, Box 8127, St. Louis, MO
          63110, USA
COMMENT
          Another IGF-I mRNA of rat liver differing in the 5' UT-region is
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ACCESSION
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REFERENCE
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           Shimatsu, A. and Rotwein, P.
 AUTHORS
            Sequence of two rat insulin-like growth factor I mRNAs differing
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  JOURNAL
           Nucleic Acids Res. 15 (17), 7196 (1987)
  MEDLINE
           88015572
           3658684
   PUBMED
REFERENCE
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 AUTHORS
           Rotwein, P.
            Direct Submission
  TITLE
            Submitted (21-OCT-1987) Rotwein P., Washington University, School
  JOURNAL
            of Medicine, 660 South Euclid Avenue, Box 8127, St. Louis, MO
            63110, USA
            Another IGF-I mRNA of rat liver differing in the 5' UT-region is
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	mRNA, clone pRIGF-1-42. Draft entry and computer-readable copy of sequence in [Mol. Endocrinol. (1987) In press] kindly

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provided by S.R.Lasky, 16-MAR-1987.
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RESULT 11
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LOCUS
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DEFINITION
ACCESSION
           AX147744
VERSION
           AX147744.1 GI:14346789
KEYWORDS
SOURCE
           Rattus norvegicus (Norway rat)
 ORGANISM
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REFERENCE
           1
 AUTHORS
           Goldspink, G.R. and Johnson, I.R.
           Use of the insulin-like-growth factor i isoform mgf for the
 TITLE
           treatment of neurological disorders
           Patent: WO 0136483-A 3 25-MAY-2001;
  JOURNAL
           University College London (GB)
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RESULT 12
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DEFINITION
         Sequence 3 from Patent W00185781.
ACCESSION
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         AX300781.1 GI:17382062
VERSION
KEYWORDS
SOURCE
         Rattus sp.
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         Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
         Rattus.
REFERENCE
 AUTHORS
         Goldspink, G.D. and Terenghi, G.B.
 TITLE
         Repair of nerve damage
         Patent: WO 0185781-A 3 15-NOV-2001;
 JOURNAL
         University College London (GB); East Grinstead Medical Research
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RESULT 13 MMIGFIBR

651 bp mRNA linear ROD 21-MAR-1995 LOCUS MMIGFIBR

Mouse mRNA for preproinsulin-like growth factor IB. DEFINITION

X04482 ACCESSION

VERSION X04482.1 GI:51806

growth factor; insulin-like growth factor IB; preproinsulin-like KEYWORDS

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growth factor IB; signal peptide.
           Mus musculus (house mouse)
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           Mus musculus
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           Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
              (bases 1 to 651)
REFERENCE
 AUTHORS
           Bell, G.I., Stempien, M.M., Fong, N.M. and Rall, L.B.
            Sequences of liver cDNAs encoding two different mouse insulin-like
 TITLE
           growth factor I precursors
           Nucleic Acids Res. 14 (20), 7873-7882 (1986)
  JOURNAL
  MEDLINE
            87040760
   PUBMED
            3774549
COMMENT
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            except for the presence of a 52 bp insertion following codon 86
            (position 397 to 448), caused by alternative RNA splicing. The B
            domain of IGF comprises residues 1-29 (position 139-225), the C
            domain residues 30-41 (position 226-261), the A domain residues
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ACCESSION VERSION	AF022961 AF022961.1 GI:2522201
KEYWORDS	
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REFERENCE	1 (bases 1 to 432)
AUTHORS TITLE	Flekna,G., Brem,G. and Mueller,M. Direct Submission
JOURNAI	Submitted (05-SEP-1997) Institute of Animal Breeding and Genetics,
	Veterinary University of Vienna, Veterinaerplatz 1, Vienna A-1210,
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         AX375028.1 GI:19169860
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 ORGANISM
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         Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
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REFERENCE
 AUTHORS
         Baak, J. and Mutter, G.L.
 TITLE
         Prognostic classification of breast cancer
         Patent: WO 0210436-A 31 07-FEB-2002;
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         THE BRIGHAM AND WOMEN'S HOSPITAL, INC. (US); Baak, Jan (US)
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Search completed: December 13, 2003, 09:27:34 Job time: 2336.77 secs

GenCore version 5.1.6 Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: December 13, 2003, 02:35:18; Search time 209.995 Seconds

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Perfect score: 523

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Searched: 2552756 seqs, 1349719017 residues

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and is derived by analysis of the total score distribution.

SUMMARIES

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	5	467.4	89.4	517	24	AAS16877	Human mechano-grow
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     10-AUG-2001 (first entry)
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KW
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PT
     Use of mechano-growth factor, an isoform of Insulin-like Growth
PT
     Factor-I, capable of reducing motoneurone loss, in the manufacture of a
PТ
     medicament for the treatment of neurological disorder -
XX
PS
     Claim 4; Page 53-54; 66pp; English.
XX
CC
     The present invention relates to use of mechano-growth factor (MGF),
CC
     an Insulin-like Growth Factor-I (IGF-I) isoform in the manufacture of a
CC
     medicament for the treatment of neurological disorder. The MGF is capable
CC
     of reducing motoneurone loss by 20% or greater in response to nerve
CC
     avulsion, and effects motoneurone rescue, preferably adult motoneurone
CC
     rescue. The MGF polynucleotide and polypeptide are useful in the
CC
     manufacture of a medicament for the treatment of a neurological disorder,
     including a disorder of motoneurones and/or neurodegenerative disorder,
CC
```

```
CC
   e.g., amyotrophic lateral sclerosis, spinal muscular atrophy, progressive
CC
   spinal muscular atrophy, infantile or juvenile muscular atrophy,
   poliomyelitis or post-polio syndrome, a disorder caused by exposure to a
CC
CC
   toxin, motoneurone trauma, a motoneurone lesion or nerve damage, an
CC
   injury that affects motoneurones, motoneurone loss associated with aging,
   autosomal or sex-linked muscular dystrophy, diabetic neuropathy,
CC
   peripheral neuropathies, Alzheimer's disease and Parkinson's disease.
CC
   The present sequence is rabbit IGF-I isoform MGF cDNA. MGF is a muscle
CC
   isoform having extracellular (Ec) domain, hence also referred as
CC
   IGF-I-Ec. The MGF protein comprises amino acid sequences encoded by
CC
CC
   nucleic acid sequence of IGF-I exons 4, 5 and 6 in the reading frame
CC
   of MGF.
XX
SO
   Sequence 523 BP; 154 A; 129 C; 142 G; 98 T; 0 other;
                    100.0%; Score 523; DB 22;
                                          Length 523;
 Query Match
                    100.0%; Pred. No. 5.1e-144;
 Best Local Similarity
 Matches 523; Conservative
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                                                     Gaps
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Db
        61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGCCACCTCAG 120
QУ
           61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
Db
       121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Qу
           121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Db
       Qу
           Db
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Qy
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Qy
           301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
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Qу
           361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Db
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Qу
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Db
       481 CAAAGATGGCATTTCCCCCAATGAAATACACAAGTAAACATTC 523
Qу
           481 CAAAGATGGCATTTCCCCCAATGAAATACACAAGTAAACATTC 523
Db
```

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     AAS16879 standard; cDNA; 523 BP.
XX
AC
     AAS16879;
XX
DT
     25-FEB-2002
                 (first entry)
XX
DE
     Rabbit mechano-growth factor (MGF) cDNA.
XX
KW
     Rabbit; mechano-growth factor; insulin-like growth factor I; IGF-I; MGF;
KW
     neuroprotective; nerve damage; peripheral nervous system; nerve severing;
KW
     muscle; neurological disorder; motoneuron loss; motorneuron disorder; ss;
KW
     nerve avulsion.
XX
     Oryctolagus cuniculus.
OS
XX
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FT
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FT
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     exon
FT
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FT
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FT
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                     /*tag= d
FT
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FT
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     exon
FT
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                     /number= 6
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PN
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PD
     15-NOV-2001.
XX
     10-MAY-2001; 2001WO-GB02054.
PF
XX
PR
     10-MAY-2000; 2000GB-0011278.
XX
PΑ
     (UNLO ) UNIV COLLEGE LONDON.
     (EGRI-) EAST GRINSTEAD MEDICAL RES TRUST.
PΑ
XX
PΙ
     Goldspink G, Terenghi G;
XX
     WPI; 2002-055585/07.
DR
     P-PSDB; AAU10561.
DR
XX
PT
     Use of insulin-like growth factor I (IGF-I) isoform known as
PT
     mechano-growth factor which is encoded by IGF-I exons 4,5,6 and has
     ability to reduce motoneuron loss in response to nerve avulsion, to
PT
PT
     treat nerve damage -
XX
PS
     Disclosure; Fig 7; 65pp; English.
```

```
XX
CC
    The invention relates to the use of an insulin-like growth factor I
CC
    (IGF-I) isoform, known as mechano-growth factor (MGF), in the manufacture
CC
    of a medicament for treating nerve damage in the peripheral nervous
CC
    system, or for treating nerve damage by localising MGF at the site of
    damage. The nerve damage may include severing of a nerve. The treatment
CC
    may be combined with another treatment (such as a polypeptide growth
CC
CC
    factor other than MGF) that prevents or diminishes degeneration of the
    target organ (for example, muscle) which the damaged nerve innervates,
CC
    whereby the treatment of the muscle with MGF or a polynucleotide encoding
CC
    MGF prevents or diminishes degeneration. The method is useful for
CC
    treating neurological disorders, preferably motorneuron disorders. These
CC
CC
    methods can reduce motoneuron loss by 20% or greater in response to nerve
    avulsion. This sequence represents cDNA encoding the rabbit MGF.
CC
XX
    Sequence 523 BP; 154 A; 129 C; 142 G; 98 T; 0 other;
SO
 Query Match
                     100.0%; Score 523; DB 24;
                    100.0%;
 Best Local Similarity
                            Pred. No. 5.1e-144;
 Matches 523; Conservative
                          0; Mismatches
                                            Indels
                                                       Gaps
                                                              0;
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Qу
           1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGGAGAC 60
Db
         61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
Qу
           61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
Db
        121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Qу
           121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Db
        Qу
           Db
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        241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
           Db
        241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
        301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGGAGTGCAGGAAACAAGAACTA 360
Qу
           301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGGAGTGCAGGAAACAAGAACTA 360
Db
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Qу
           361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Db
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QУ
           421 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTT 480
Db
        481 CAAAGATGGCATTTCCCCCCAATGAAATACACAAGTAAACATTC 523
Qу
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481 CAAAGATGGCATTTCCCCCCAATGAAATACACAAGTAAACATTC 523

Db

```
RESULT 3
AAT84893
     AAT84893 standard; cDNA; 553 BP.
XX
AC
     AAT84893;
XX
DT
     14-APR-1998 (first entry)
XX
     Rabbit insulin like growth factor 1 encoding cDNA.
DE
XX
     Insulin like growth factor 1; IGF-1; Ec peptide; muscle disorder;
KW
KW
     heart; neuromuscular disease; primer; ss.
XX
     Oryctolagus cuniculus.
OS
XX
                     Location/Qualifiers
FH
                     1..366
FT
     CDS
                     /*tag= a
FT
                     /product= "IGF-1"
FT
XX.
PN
     WO9733997-A1.
XX
PD
     18-SEP-1997.
XX
                    97WO-GB00658.
PF
     11-MAR-1997;
XX
     11-MAR-1996;
                    96GB-0005124.
PR
XX
     (UNLO ) ROYAL FREE HOSPITAL SCHOOL MED.
PA
XX
     Goldspink G;
PΙ
ХX
     WPI; 1997-470877/43.
DR
     P-PSDB; AAW23301.
DR
ХХ
     Use of insulin like growth factor I characterised by presence of Ec
PT
     peptide - to treat humans or animals, particularly muscle disorders,
PT
     heart conditions or neuromuscular diseases
PT
XX
     Disclosure; Fig 3; 33pp; English.
PS
XX
     A use of insulin like growth factor I (IGF-1) has been developed, and
CC
     is characterised by the presence of the Ec peptide, or a functional
CC
     equivalent, in the treatment or therapy of a human or animal. The IGF-1
CC
     polypeptide can be used to treat muscular disorders, e.g. Duchenne or
CC
     Becker muscular dystrophy, autosomal dystrophies and related progressive
CC
     skeletal muscle weakness and wasting, muscle atrophy in ageing humans,
CC
     spinal cord injury induced muscle atrophy and neuromuscular diseases,
CC
     and cardiac disorders, e.g. diseases where promotion of cardiac muscle
CC
     protein synthesis is a beneficial treatment, cardiomyopathies and acute
CC
     heart failure or insult, specifically myocarditis or myocardial
CC
     infarction. It can also be used to promote bone fracture healing and
CC
     maintenance of bone in old age. The present sequence encodes rabbit
CC
CC
     IGF-1 used in the present specification.
XX
     Sequence 553 BP; 159 A; 142 C; 147 G; 105 T; 0 other;
SQ
```

```
Query Match
                         Score 523; DB 18; Length 553;
                   100.0%;
 Best Local Similarity
                  100.0%; Pred. No. 5.3e-144;
 Matches 523; Conservative
                       0; Mismatches
                                     0;
                                        Indels
                                                  Gaps
                                                        0;
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Qy
          31 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGGAGAC 90
Dh
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Qу
          Db
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       121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Qу
          Db
       151 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 210
       Qу
          Db
       241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
Qу
          271 ATGCCCAAGACTCCAGAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 330
Db
       301 AGAAGGAAAGGAAGTACATTTGAAGAACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
Qу
          331 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 390
Db
       361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Qу
          391 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 450
Db
       421 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTT 480
Qу
          Db
       451 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTT 510
       481 CAAAGATGGCATTTCCCCCCAATGAAATACACAAGTAAACATTC 523
Qу
           Db
       511 CAAAGATGGCATTTCCCCCCAATGAAATACACAAGTAAACATTC 553
RESULT 4
TD
   AAD06398 standard; cDNA; 517 BP.
XX
AC
   AAD06398;
XX
DT
   10-AUG-2001 (first entry)
XX
   Human IGF-I isoform mechano-growth factor (MGF) cDNA.
DE
XX
   Human; IGF-I isoform; Insulin-like Growth Factor-I; MGF;
KW
KW
   mechano-growth factor; neurological disorder; neurodegenerative disorder;
   amyotrophic lateral sclerosis; spinal muscular atrophy; muscular atrophy;
KW
KW
   poliomyelitis; post-polio syndrome; toxin; motoneurone disorder;
```

nerve damage; autosomal muscular dystrophy; diabetic neuropathy;

KW

```
KW
     sex-linked muscular dystrophy; peripheral neuropathy;
KW
     Alzheimer's disease; Parkinson's disease; ss.
XX
OS
     Homo sapiens.
XX
FΗ
                     Location/Qualifiers
     Kev
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     CDS
                     1..333
FT
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                     /note= "This region comprises exons 3-6. The CDS does
FT
FΤ
                     not include start codon"
                     /partial
FT
XX
PN
     WO200136483-A1.
XX
PΩ
     25-MAY-2001.
XX
ΡF
     15-NOV-2000; 2000WO-GB04354.
XX
PR
     15-NOV-1999;
                    99GB-0026968.
XX
     (UNLO ) UNIV COLLEGE LONDON.
PΑ
XX
     Goldspink G, Johnson I;
PI
XX
DR
     WPI; 2001-355620/37.
DR
     P-PSDB; AAE02447.
XX
     Use of mechano-growth factor, an isoform of Insulin-like Growth
PT
PT
     Factor-I, capable of reducing motoneurone loss, in the manufacture of a
     medicament for the treatment of neurological disorder -
PT
XX
PS
     Claim 4; Page 49-50; 66pp; English.
XX
     The present invention relates to use of mechano-growth factor (MGF),
CC
CC
     an Insulin-like Growth Factor-I (IGF-I) isoform in the manufacture of a
     medicament for the treatment of neurological disorder. The MGF is capable
CC
     of reducing motoneurone loss by 20% or greater in response to nerve
CC
     avulsion, and effects motoneurone rescue, preferably adult motoneurone
CC
     rescue. The MGF polynucleotide and polypeptide are useful in the
CC
CC
     manufacture of a medicament for the treatment of a neurological disorder,
     including a disorder of motoneurones and/or neurodegenerative disorder,
CC
     e.g., amyotrophic lateral sclerosis, spinal muscular atrophy, progressive
CC
CC
     spinal muscular atrophy, infantile or juvenile muscular atrophy,
     poliomyelitis or post-polio syndrome, a disorder caused by exposure to a
CC
     toxin, motoneurone trauma, a motoneurone lesion or nerve damage, an
CC
CC
     injury that affects motoneurones, motoneurone loss associated with aging,
CC
     autosomal or sex-linked muscular dystrophy, diabetic neuropathy,
     peripheral neuropathies, Alzheimer's disease and Parkinson's disease.
CC
CC
     The present sequence is human IGF-I isoform MGF cDNA. MGF is a muscle
     isoform having extracellular (Ec) domain, hence also referred as
CC
CC
     IGF-I-Ec. The MGF protein comprises amino acid sequences encoded by
CC
     nucleic acid sequence of IGF-I exons 4, 5 and 6 in the reading frame
CC
     of MGF.
XX
     Sequence 517 BP; 150 A; 130 C; 139 G; 98 T; 0 other;
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SO

```
89.4%; Score 467.4; DB 22; Length 517;
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 Best Local Similarity
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                                                4; Gaps
                                                          2;
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Db
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           61 AGGGGCTTTTATTTCAACAAGCCCACAGGGTATGGCTCCAGCAGTCGGAGGGCGCCTCAG 120
Dh
       121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Qy
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Db
       Qу
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Db
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Qу
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Db
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Qу
           298 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 357
Db
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Qу
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Db
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Qу
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Db
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Qу
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Db
RESULT 5
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TD
XX
AC
    AAS16877;
ХX
DT
    25-FEB-2002 (first entry)
XX
DE
    Human mechano-growth factor (MGF) cDNA.
ХX
    Human; mechano-growth factor; insulin-like growth factor I; IGF-I; MGF;
KW
    neuroprotective; nerve damage; peripheral nervous system; nerve severing;
KW
    muscle; neurological disorder; motoneuron loss; motorneuron disorder; ss;
KW
    nerve avulsion.
KW
XX
```

OS

Homo sapiens.

```
XX
FΗ
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PR
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PΑ
     (EGRI-) EAST GRINSTEAD MEDICAL RES TRUST.
PA
XX
     Goldspink G,
                   Terenghi G;
PΙ
XX
     WPI; 2002-055585/07.
DR
DR
     P-PSDB; AAU10559.
XX
PT
     Use of insulin-like growth factor I (IGF-I) isoform known as
РΤ
     mechano-growth factor which is encoded by IGF-I exons 4,5,6 and has
PΤ
     ability to reduce motoneuron loss in response to nerve avulsion, to
PT
     treat nerve damage -
XX
PS
     Claim 11; Fig 5; 65pp; English.
XX
CC
     The invention relates to the use of an insulin-like growth factor I
CC
     (IGF-I) isoform, known as mechano-growth factor (MGF), in the manufacture
CC
     of a medicament for treating nerve damage in the peripheral nervous
CC
     system, or for treating nerve damage by localising MGF at the site of
CC
     damage. The nerve damage may include severing of a nerve. The treatment
CC
     may be combined with another treatment (such as a polypeptide growth
     factor other than MGF) that prevents or diminishes degeneration of the
CC
CC
     target organ (for example, muscle) which the damaged nerve innervates,
CC
     whereby the treatment of the muscle with MGF or a polynucleotide encoding
CC
     MGF prevents or diminishes degeneration. The method is useful for
     treating neurological disorders, preferably motorneuron disorders. These
CC
CC
     methods can reduce motoneuron loss by 20% or greater in response to nerve
CC
     avulsion. This sequence represents cDNA encoding the human MGF.
XX
```

```
89.4%; Score 467.4; DB 24; Length 517;
 Query Match
 Best Local Similarity
                   96.2%; Pred. No. 1.3e-127;
                           Mismatches
                                      16;
 Matches 501; Conservative
                                                          2;
                         0;
                                         Indels
                                                 4; Gaps
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Qу
          1 GGACCGGAGACGCTCTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGGAGAC 60
Db
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          61 AGGGGCTTTTATTTCAACAAGCCCACAGGGTATGGCTCCAGCAGTCGGAGGGCGCCTCAG 120
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Db
       QУ
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Db
       241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
QУ
          241 ATGCCCAAGACCCAGAAGTATCAGCCCCCATCTACCAACAAGAACACGAAGTCTCA---G 297
Db
       301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
Qу
          298 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 357
Db
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Qу
           358 CAGGATGTA-GAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 416
Db
       421 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTT 480
Qу
           417 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTT 476
Db
       481 CAAAGATGGCATTTCCCCCAATGAAATACACAAGTAAACAT 521
Qу
           477 CAAAGATGGCATTTCCCCCAATGAAATACACAAGTAAACAT 517
Dh
RESULT 6
AAD06405
ID
   AAD06405 standard; cDNA; 471 BP.
ХX
AC
   AAD06405;
XX
DT
    10-AUG-2001 (first entry)
XX
DE
    Rabbit liver-type IGF-I isoform (L.IGF-I) cDNA.
XX
    Rabbit; IGF-I isoform; Insulin-like Growth Factor-I; MGF;
KW
    mechano-growth factor; neurological disorder; neurodegenerative disorder;
KW
KW
    amyotrophic lateral sclerosis; spinal muscular atrophy; muscular atrophy;
KW
   poliomyelitis; post-polio syndrome; toxin; motoneurone disorder;
```

```
nerve damage; autosomal muscular dystrophy; diabetic neuropathy;
KW
     sex-linked muscular dystrophy; peripheral neuropathy;
KW
     Alzheimer's disease; Parkinson's disease; liver; L.IGF-I; ss.
KW
XX
     Oryctolagus cuniculus.
OS
XX
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FΗ
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FT
                     /transl except= (pos:25..27, aa:Gln)
FT
                     /note= "These translation exceptions occur while decoding
FT
                     the alternative version of the protein (AAE02456).
FT
                     The CDS comprises exons 3, 4 and 6 and
FT
                     does not include start codon"
FT
FT
                     /partial
XX
PN
     WO200136483-A1.
XX
     25-MAY-2001.
PD:
XX
     15-NOV-2000; 2000WO-GB04354.
PF
XX
PR
     15-NOV-1999:
                    99GB-0026968.
XX
     (UNLO ) UNIV COLLEGE LONDON.
PΑ
XX
PΙ
     Goldspink G, Johnson I;
XX
DR -
     WPI; 2001-355620/37.
DR
     P-PSDB; AAE02452, AAE02456.
ХX
PT
     Use of mechano-growth factor, an isoform of Insulin-like Growth
     Factor-I, capable of reducing motoneurone loss, in the manufacture of a
PT
     medicament for the treatment of neurological disorder -
PT
XX
PS
     Disclosure; Page 59-60; 66pp; English.
XX
     The present invention relates to use of mechano-growth factor (MGF),
CC
     an Insulin-like Growth Factor-I (IGF-I) isoform in the manufacture of a
CC
     medicament for the treatment of neurological disorder. The MGF is capable
CC
     of reducing motoneurone loss by 20% or greater in response to nerve
CC
CC
     avulsion, and effects motoneurone rescue, preferably adult motoneurone
     rescue. The MGF polynucleotide and polypeptide are useful in the
CC
     manufacture of a medicament for the treatment of a neurological disorder,
CC
     including a disorder of motoneurones and/or neurodegenerative disorder,
CC
     e.g., amyotrophic lateral sclerosis, spinal muscular atrophy, progressive
CC
CC
     spinal muscular atrophy, infantile or juvenile muscular atrophy,
CC
     poliomyelitis or post-polio syndrome, a disorder caused by exposure to a
CC
     toxin, motoneurone trauma, a motoneurone lesion or nerve damage, an
     injury that affects motoneurones, motoneurone loss associated with aging,
CC
     autosomal or sex-linked muscular dystrophy, diabetic neuropathy,
CC
     peripheral neuropathies, Alzheimer's disease and Parkinson's disease.
CC
CC
     The present sequence is rabbit liver-type IGF-I isoform (L.IGF-I) cDNA.
     The L.IGF-I protein comprises amino acid sequences encoded by
CC
CC
     nucleic acid sequence of IGF-I exons 4 and 6.
```

```
XX
```

SQ Sequence 471 BP; 132 A; 118 C; 131 G; 90 T; 0 other;

```
78.2%; Score 409; DB 22; Length 471;
 Query Match
 Best Local Similarity
                  90.1%; Pred. No. 2e-110;
 Matches 471; Conservative
                       0; Mismatches
                                    0; Indels
                                                       1;
        1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGAGAC 60
Qу
          1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 60
Db
        61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
Qу
          Db
       121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Qу
          121 ACAGGCATCGTGGATGATGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Db
       Qу
          Db
       241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
Qу
          241 ATGCCCAAGACTCAG--
Db
       301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
Qу
               256 -----AAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 308
Db
       361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Qу
          309 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 368
Db
       421 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTT 480
Qу
          Db
       369 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTT 428
       481 CAAAGATGGCATTTCCCCCAATGAAATACACAAGTAAACATTC 523
Qу
          Db
       429 CAAAGATGGCATTTCCCCCAATGAAATACACAAGTAAACATTC 471
RESULT 7
TD
   AAS16884 standard; cDNA; 471 BP.
XX
AC
   AAS16884;
XX
DT
   25-FEB-2002 (first entry)
XX
DE
   Rabbit insulin-like growth factor I liver-type isoform (L.IGF-I) cDNA.
XX
KW
   Rabbit; mechano-growth factor; insulin-like growth factor I; IGF-I; MGF;
KW
   neuroprotective; nerve damage; peripheral nervous system; nerve severing;
KW
   muscle; neurological disorder; motoneuron loss; motorneuron disorder; ss;
```

```
nerve avulsion; insulin-like growth factor I liver-type isoform; L.IGF-I;
KW
XX
OS
     Oryctolagus cuniculus.
XX
                     Location/Qualifiers
FΗ
     Key
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FT
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                     /*taq=a
FT
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FT
FT
                     /partial
                     /note= "No start codon"
FT
FT
                     1..75
     exon
FT
                     /*taq=b
FT
                     /number= exon 3
                     76..258
FT
     exon
                     /*tag= c
FT
                     /number= exon 4
FT
                     259..315
FT
     exon
                     /*tag= d
FT
FT
                     /number= exon 6
XX
PN
     WO200185781-A2.
ХХ
PD
     15-NOV-2001.
XX
PF
     10-MAY-2001; 2001WO-GB02054.
XX
PR
     10-MAY-2000; 2000GB-0011278.
XX
PΑ
     (UNLO ) UNIV COLLEGE LONDON.
     (EGRI-) EAST GRINSTEAD MEDICAL RES TRUST.
PΑ
XX
PΙ
                   Terenghi G;
     Goldspink G,
XX
DR
     WPI; 2002-055585/07.
DR
     P-PSDB; AAU10564.
XX
     Use of insulin-like growth factor I (IGF-I) isoform known as
PT
PT
     mechano-growth factor which is encoded by IGF-I exons 4,5,6 and has
     ability to reduce motoneuron loss in response to nerve avulsion, to
PT
PT
     treat nerve damage -
XX
PS
     Disclosure; Fig 10; 65pp; English.
XX
CC
     The invention relates to the use of an insulin-like growth factor I
CC
     (IGF-I) isoform, known as mechano-growth factor (MGF), in the manufacture
CC
     of a medicament for treating nerve damage in the peripheral nervous
CC
     system, or for treating nerve damage by localising MGF at the site of
CC
     damage. The nerve damage may include severing of a nerve. The treatment
CC
     may be combined with another treatment (such as a polypeptide growth
CC
     factor other than MGF) that prevents or diminishes degeneration of the
CC
     target organ (for example, muscle) which the damaged nerve innervates,
CC
     whereby the treatment of the muscle with MGF or a polynucleotide encoding
     MGF prevents or diminishes degeneration. The method is useful for
CC
     treating neurological disorders, preferably motorneuron disorders. These
CC
     methods can reduce motoneuron loss by 20% or greater in response to nerve
CC
CC
     avulsion. This sequence represents cDNA encoding the rabbit insulin-like
     growth factor I liver-type isoform (L.IGF-I) used in experiments on
```

```
CC
   motoneuron loss.
XX
   Sequence 471 BP; 132 A; 118 C; 131 G; 90 T; 0 other;
SO
 Query Match
                   78.2%; Score 409; DB 24; Length 471;
 Best Local Similarity
                   90.1%; Pred. No. 2e-110;
                        0; Mismatches
                                     0; Indels
                                                        1;
 Matches 471; Conservative
                                               52:
                                                  Gaps
        1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGGAGAC 60
Qу
          1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGAGATGCTCTTCAGTTCGTGTGTGGAGAC 60
Db
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Qу
          Db
        61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
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Qу
          121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Db
       Qу
          Db
       241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
Qу
          241 ATGCCCAAGACTCAG----
Db
       301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
Qу
               256 -----AAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 308
Db
       361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Qу
          309 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 368
Db
       421 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTT 480
Qy
           369 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTT 428
Db
       481 CAAAGATGGCATTTCCCCCAATGAAATACACAAGTAAACATTC 523
Qу
          429 CAAAGATGGCATTTCCCCCAATGAAATACACAAGTAAACATTC 471
Db
RESULT 8
AAD06399
    AAD06399 standard; cDNA; 539 BP.
ID
XX
AC
    AAD06399;
XX
DT
    10-AUG-2001 (first entry)
XX
    Rat IGF-I isoform mechano-growth factor (MGF) cDNA.
DE
XX
    Rat; IGF-I isoform; Insulin-like Growth Factor-I; MGF;
KW
    mechano-growth factor; neurological disorder; neurodegenerative disorder;
KW
```

```
amyotrophic lateral sclerosis; spinal muscular atrophy; muscular atrophy;
KW
     poliomyelitis; post-polio syndrome; toxin; motoneurone disorder;
KW
     nerve damage; autosomal muscular dystrophy; diabetic neuropathy;
KW
     sex-linked muscular dystrophy; peripheral neuropathy;
KW
     Alzheimer's disease; Parkinson's disease; ss.
KW
XX
OS
     Rattus sp.
XX
                     Location/Qualifiers
FH
     Key
FΤ
     CDS
                     1..336
FT
                     /*taq=a
                     /product= "Mechano-growth factor (MGF)"
FT ·
                     /note= "This region comprises exons 3-6. The CDS does
FT
                     not include start codon"
FT
                     /partial
FT
XX
PN
     WO200136483-A1.
XX
PD
     25-MAY-2001.
XX
PF
     15-NOV-2000; 2000WO-GB04354.
XX
PR
                    99GB-0026968.
     15-NOV-1999;
XX
PΑ
     (UNLO ) UNIV COLLEGE LONDON.
XX
PΙ
     Goldspink G, Johnson I;
XX
DR
     WPI; 2001-355620/37.
DR
     P-PSDB; AAE02448.
XX
PT
     Use of mechano-growth factor, an isoform of Insulin-like Growth
     Factor-I, capable of reducing motoneurone loss, in the manufacture of a
PТ
     medicament for the treatment of neurological disorder -
PΤ
XX
     Claim 4; Page 51-52; 66pp; English.
PS
XX
     The present invention relates to use of mechano-growth factor (MGF),
CC
     an Insulin-like Growth Factor-I (IGF-I) isoform in the manufacture of a
CC
     medicament for the treatment of neurological disorder. The MGF is capable
CC
     of reducing motoneurone loss by 20% or greater in response to nerve
CC
     avulsion, and effects motoneurone rescue, preferably adult motoneurone
CC
CC
     rescue. The MGF polynucleotide and polypeptide are useful in the
     manufacture of a medicament for the treatment of a neurological disorder,
CC
     including a disorder of motoneurones and/or neurodegenerative disorder,
CC
     e.q., amyotrophic lateral sclerosis, spinal muscular atrophy, progressive
CC
     spinal muscular atrophy, infantile or juvenile muscular atrophy,
CC
     poliomyelitis or post-polio syndrome, a disorder caused by exposure to a
CC
     toxin, motoneurone trauma, a motoneurone lesion or nerve damage, an
CC
     injury that affects motoneurones, motoneurone loss associated with aging,
CC
     autosomal or sex-linked muscular dystrophy, diabetic neuropathy,
CC
CC
     peripheral neuropathies, Alzheimer's disease and Parkinson's disease.
     The present sequence is rat IGF-I isoform MGF cDNA. MGF is a muscle
CC
CC
     isoform having extracellular (Ec) domain, hence also referred as
CC
     IGF-I-Ec. The MGF protein comprises amino acid sequences encoded by
     nucleic acid sequence of IGF-I exons 4, 5 and 6 in the reading frame
CC
CC
     of MGF.
```

```
Sequence 539 BP; 161 A; 136 C; 139 G; 103 T; 0 other;
SO
 Query Match
                    68.2%;
                          Score 356.8; DB 22; Length 539;
 Best Local Similarity
                    82.3%;
                          Pred. No. 5.1e-95;
 Matches 436; Conservative
                         0; Mismatches
                                      87;
                                          Indels
                                                  7;
                                                     Gaps
                                                           2;
         1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGGAGAC 60
Qу
           Db
         1 GGACCAGAGACCCTTTGCGGGGCTGAGCTGGTGGACGCTCTTCAGTTCGTGTGTGGACCA 60
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Qу
           Db
        61 AGGGGCTTTTACTTCAACAAGCCCACAGTCTATGGCTCCAGCATTCGGAGGGCACCACAG 120
Qy
        121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
           Db
        121 ACGGCATTGTGGATGAGTGTTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
        Qу
              Db
        181 TGTGTCCGCTGCAAGCCTACAAAGTCAGCTCGTTCCATCCGGGCCCAGCGCCACACTGAC 240
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           Db
        241 ATGCCCAAGACTCAGAAGTCCCAGCCCCTATCGACACACAAGAAAAGGAAGCTGCAAAGG 300
       301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGGAGTGCAGGAAACAAGAACTA 360
QУ
           301 AGAAGGAAAGGAAGTACACTTGAAGAACACAAGTAGAGGAAGTGCAGGAAACAAGACCTA 360
Db
       361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
QУ
           Db
        361 CAGAATGTAGGAGGAGCCTCCCGAGGAACAGAAAATGCCACGTCACCGCAAGATCCTTTG 420
        421 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCA----AAAAATAAGTTTGATC 474
Ov
                         Db
       421 CTGCTTGAGCAACCTGCAAAACATCGGAACACCTGCCAAATATCAATAATGAGTTCAATA 480
Qу
        475 ACATTTCAAAGAT-GGCATTTCCCCCAATGAAATACACAAGTAAACATTC 523
            Db
       481 TCATTTCAGAGATGGGCATTTCCCTCAATGAAATACACAAGTAAACATTC 530
RESULT 9
    AAS16878 standard; cDNA; 539 BP.
XX
AC
   AAS16878;
XX
DT
    25-FEB-2002 (first entry)
XX
DE
   Rat mechano-growth factor (MGF) cDNA.
XX
    Rat; mechano-growth factor; insulin-like growth factor I; IGF-I; MGF;
KW
KW
    neuroprotective; nerve damage; peripheral nervous system; nerve severing;
    muscle; neurological disorder; motoneuron loss; motorneuron disorder; ss;
KW
```

XX

```
KW
    nerve avulsion.
XX
OS
    Rattus sp.
ΧX
FΗ
                     Location/Qualifiers
     Key
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                     1..336
     CDS
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FT
FΤ
                     /partial
                     /note= "No start codon"
FT
FT
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FT
                     76..258
     exon
FT
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FT
                     259..309
FT
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                     /*tag= d
FT
                     /number= exon 5
FT
                     310..333
FT
     exon
                     /*tag=e
FT
FT
                     /number= exon 6
XX
PN
     WO200185781-A2.
XX
     15-NOV-2001.
PD
XX
     10-MAY-2001; 2001WO-GB02054.
PF
XX
     10-MAY-2000; 2000GB-0011278.
PR
XX
     (UNLO ) UNIV COLLEGE LONDON.
PA
PΑ
     (EGRI-) EAST GRINSTEAD MEDICAL RES TRUST.
XX
PΙ
     Goldspink G, Terenghi G;
XX
     WPI; 2002-055585/07.
DR
DR
     P-PSDB; AAU10560.
XX
     Use of insulin-like growth factor I (IGF-I) isoform known as
PT
PΤ
     mechano-growth factor which is encoded by IGF-I exons 4,5,6 and has
PT
     ability to reduce motoneuron loss in response to nerve avulsion, to
PT
     treat nerve damage -
XX
PS
     Disclosure; Fig 6; 65pp; English.
XX
     The invention relates to the use of an insulin-like growth factor I
CC
     (IGF-I) isoform, known as mechano-growth factor (MGF), in the manufacture
CC
     of a medicament for treating nerve damage in the peripheral nervous
CC
     system, or for treating nerve damage by localising MGF at the site of
CC
     damage. The nerve damage may include severing of a nerve. The treatment
CC
     may be combined with another treatment (such as a polypeptide growth
CC
CC
     factor other than MGF) that prevents or diminishes degeneration of the
     target organ (for example, muscle) which the damaged nerve innervates,
CC
     whereby the treatment of the muscle with MGF or a polynucleotide encoding
CC
     MGF prevents or diminishes degeneration. The method is useful for
CC
     treating neurological disorders, preferably motorneuron disorders. These
CC
```

```
avulsion. This sequence represents cDNA encoding the rat MGF.
CC
XX
   Sequence 539 BP; 161 A; 136 C; 139 G; 103 T; 0 other;
SO
                   68.2%; Score 356.8; DB 24; Length 539;
 Ouery Match
 Best Local Similarity
                   82.3%; Pred. No. 5.1e-95;
                         0; Mismatches
                                      87; Indels
 Matches 436; Conservative
         1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGGAGAC 60
Qу
          1 GGACCAGAGACCCTTTGCGGGGCTGAGCTGGTGGACGCTCTTCAGTTCGTGTGTGGACCA 60
Db
        61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
Qу
           61 AGGGGCTTTTACTTCAACAAGCCCACAGTCTATGGCTCCAGCATTCGGAGGGCACCACAG 120
Db
       121 ACAGGCATCGTGGATGATGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Qу
           121 ACGGGCATTGTGGATGAGTGTTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Db
       QУ
           181 TGTGTCCGCTGCAAGCCTACAAAGTCAGCTCGTTCCATCCGGGCCCAGCGCCACACTGAC 240
Db
       241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
Qу
           241 ATGCCCAAGACTCAGAAGTCCCAGCCCCTATCGACACACAAGAAAAGGAAGCTGCAAAGG 300
Db
       301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
Qу
           301 AGAAGGAAAGGAAGTACACTTGAAGAACACAAGTAGAGGAAGTGCAGGAAACAAGACCTA 360
Db
       361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Qу
           361 CAGAATGTAGGAGGAGCCTCCCGAGGAACAGAAAATGCCACGTCACCGCAAGATCCTTTG 420
Db
       421 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCA-----AAAAATAAGTTTGATC 474
Qу
                        421 CTGCTTGAGCAACCTGCAAAACATCGGAACACCTGCCAAATATCAATAATGAGTTCAATA 480
Db
       475 ACATTTCAAAGAT-GGCATTTCCCCCAATGAAATACACAAGTAAACATTC 523
Qу
           481 TCATTTCAGAGATGGGCATTTCCCTCAATGAAATACACAAGTAAACATTC 530
Db
RESULT 10
ABV76185
    ABV76185 standard; cDNA; 651 BP.
XX
AC
    ABV76185;
XX
DT
    07-MAR-2003 (first entry)
XX
    Mouse insulin-like growth factor IB cDNA.
DE
XX
KW
    Insulin-like growth factor IB; IGF-IB; mouse; mRNA; assay;
```

methods can reduce motoneuron loss by 20% or greater in response to nerve

CC

```
KW
     nucleic acid detection; gene; ss.
XX
OS
     Mus musculus.
XX
FΗ
                     Location/Qualifiers
     Key
                     73..474
FT
     CDS
FT
                     /*taq=a
                     /product= "IGF-IB"
FT
XX
     WO200297390-A2.
PN
XX
PD
     05-DEC-2002.
XX
PF
     31-MAY-2002; 2002WO-SE01056.
XX
     01-JUN-2001; 2001SE-0001934.
PR
XX
PA
     (BIOV-) BIOVITRUM AB.
XX
PΙ
     Parrow V, Rosengren L;
XX
DR
     WPI; 2003-129529/12.
XX
     Quantitating a target nucleic acid in a sample comprises immobilizing,
PΤ
     on a solid support, a sample comprising a target nucleic acid, and
PT
     detecting and quantitating signals generated from the antisense and
PT
PT
     sense probes -
XX
     Example 1; Page 16-17; 18pp; English.
PS
XX
     The present sequence is that of cDNA encoding murine insulin-like
CC
     growth factor 1B (IGF-IB). The cDNA was used in an example of the
CC
     method of the invention to generate probes for determination of
CC
CC
     IGF-IB RNA. The method comprises a quantitative hybridisation
     assay for analysis of mRNA in a target nucleic acid (TNA) sample.
CC
CC
     It involves: (i) immobilising the TNA sample on a solid support;
     (ii) contacting a labelled antisense probe to a first portion of the
CC
     TNA, and a labelled sense probe to a second portion of the TNA;
CC
     (iii) detecting and quantitating the signals generated from the
CC
CC
     hybridised probes; and (iv) determining the value represented by
CC
     the antisense probe signal minus the sense probe signal, the value
     being proportional to the amount of mRNA in the TNA sample. In an
CC
     example of the method, a cDNA clone containing 60 nucleotides from
CC
     exon 2 and 179 nucleotides from exon 3 of the mouse IGF-IB gene was
CC
CC
     cloned into pGEN-4Z vector. Linearisation of the plasmid with
CC
     EcoRI allowed transcription of a 250-nucleotide antisense probe
CC
     using T7 polymerase. Linearisation with HindIII allowed
CC
     transcription of a sense probe of similar length using SP6
     polymerase (see ABV76186). The probes were purified and used to
CC
     determine IGF-I RNA in mouse hepatocytes and also in rat hepatocytes.
CC
XX
SO
     Sequence 651 BP; 193 A; 185 C; 149 G; 124 T; 0 other;
                          66.8%; Score 349.4; DB 25; Length 651;
  Query Match
                          82.8%; Pred. No. 8.3e-93;
  Best Local Similarity
                                                   81; Indels
                                                                  7; Gaps
                                                                              2;
  Matches 425; Conservative
                                 0; Mismatches
```

```
1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGGAGAC 60
Qу
          139 GGACCAGAGACCCTTTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGGACCG 198
Db
        61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
QУ
          199 AGGGGCTTTTACTTCAACAAGCCCACAGGCTATGGCTCCAGCATTCGGAGGGCACCTCAG 258
Db
       121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
QУ
          259 ACAGGCATTGTGGATGAGTGTTGCTTCCGGAGCTGTGATCTGAGGAGACTGGAGATGTAC 318
Db
       QУ
          319 TGTGCCCCACTGAAGCCTACAAAAGCAGCCCGCTCTATCCGTGCCCAGCGCCACACTGAC 378
Db
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Qу
          Db
       301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
Oy
          439 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGAAGTGCAGGAAACAAGACCTA 498
Db
       361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Qу
                           | | | | |
                                499 CAGAATGTAGGAGGAGCCTCCCACGGAGCAGAAAATGCCACATCACCGCAGGATCCTTTG 558
Db
                                         -AAAAATAAGTTTGATC 474
       421 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCA--
Qу
                       559 CTGCTTGAGCAACCTGCAAAACATCGAAACACCTACCAAATAACAATAATAAGTCCAATA 618
Db
       475 ACATTTCAAAGAT-GGCATTTCCCCCAATGAAA 506
Qу
          Db
       619 ACATTACAAAGATGGGCATTTCCCCCAATGAAA 651
RESULT 11
AAN70436
ID
   AAN70436 standard; cDNA; 818 BP.
XX
AC
   AAN70436;
XX
DT
   25-MAR-2003
             (updated)
            (first entry)
DT
   05-APR-1991
XX
   Sequence encoding insulin-like growth factor 1A (IGF-1A).
DE
XX
KW
   Growth promoter; lactation enhancer; cell proliferation; ss.
XX
OS
   Homo sapiens.
ХХ
ΡN
   EP229750-A.
XX
PD
   22-JUL-1987.
XX
ΡF
   06-JAN-1987;
              87EP-0870001.
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XX
PR
    20-NOV-1986;
                 86US-0929671.
PR
                 86US-0816662.
    07-JAN-1986;
XX
PΑ
    (UNIW ) UNIV WASHINGTON.
ХX
    Krivi GG, Rotwein PS;
PΙ
XX
DR
    WPI: 1987-200203/29.
XX
PT
    New pre-pro-insulin-like growth factor-1 protein - obtd. by
РΤ
    recombinant DNA procedures for use as growth promoters for
РΤ
    enhancing lactation, for stimulating cell proliferation etc.
XX
PS
    Example; Fig 5; 59pp; English.
XX
CC
    A 42 base oligonucleotide corresponding to the DNA sequence encoding
CC
    amino acids 10 to 23 of mature human IGF-I was synthesized (AAN70437).
CC
    The radiolabeled 42 mer was then employed to screen for IGF-I
CC
    containing DNA sequences in a human liver cDNA library. Insulin-
CC
    like growth factors-1A and -1B cDNAs were isolated from a human cDNA
   ·library by using lambdagt 11 (AAN70435, AAN70436). The human IGF-1
CC
CC
    genomic gene was isolated and mapped. It encodes at least two
CC
    preproinsulin-like growth factor-1 proteins. An essentially pure
CC
    proproinsulin-like growth factor-1 protein comprising the sequence
CC
    of amino acids shown in Figure six is claimed (AAP70277).
CC
    (Updated on 25-MAR-2003 to correct PA field.)
XX
    Sequence 818 BP; 232 A; 186 C; 187 G; 213 T; 0 other;
SO
                      63.9%; Score 334.4; DB 8; Length 818;
 Ouery Match
 Best Local Similarity
                      84.6%; Pred. No. 2.4e-88;
                           0; Mismatches 26;
 Matches 445; Conservative
                                               Indels
                                                                  4;
          1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 60
Qу
            Db
        203 GGACCGGAGACGCTCTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 262
         61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
Qу
            263 AGGGGCTTTTATTTCAACAAGCCCACAGGGTATGGCTCCAGCAGTCGGAGGGCGCCTCAG 322
Dh
        121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Qу
            323 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTAAGGAGGCTGGAGATGTAT 382
Db
        QУ
            Db
        383 TGCGCACCCTCAAGCCTGCCAAGTCAGCTCGCTCTGTCCGTGCCCAGCGCCACACCGAC 442
        241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
QУ
            Db
        443 ATGCCCAAGACCCAG--
        301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
Qу
                  Db
        458 -----AAGGAAGTACATTTGAAGAACGCAAGTAGAGGGAGTGCAGGAAACAAGAACTA 510
```

```
361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Qу
             511 CAGGATGTAGGAAGACCCTCCTGAGGAGTGAAGAGTGACATGCCACCGCAGGATCCTTTG 570
Db
         421 CTCTGCAC-AGTTACCTG-TAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACAT 478
Oy
             571 CTCTGCACGAGTTACCTGTTAAACTTTGGAACACCTACCAAAAAATAAGTTTGATAACAT 630
Db
         479 TTCAAAGAT-GGCATTTCCCCCAATGAAATACACAAGTAAACATTC 523
Qу
             Db
         631 TTAAAAGATGGGCGTTTCCCCCAATGAAATACACAAGTAAACATTC 676
RESULT 12
ABT11091
    ABT11091 standard; cDNA; 7260 BP.
XX
AC
    ABT11091;
XX
DT
    04-DEC-2002 (first entry)
XX
DE
    Human breast cancer associated coding sequence SEQ ID NO: 1225.
XX
KW
    Human; breast specific gene; breast cancer; differential expression;
KW
    cytostatic; gene therapy; gene; ss.
XX
OS
    Homo sapiens.
XX
PN
    WO200259271-A2.
XX
PD
    01-AUG-2002.
XX
PF
    25-JAN-2002; 2002WO-US02176.
XX
PR
    25-JAN-2001; 2001US-263757P.
    25-APR-2001; 2001US-286090P.
PR
    23-MAY-2001; 2001US-292517P.
PR
XX
PΑ
    (GENE-) GENE LOGIC INC.
XX
ΡI
    Orr MS, Nation M, Diggans JC, Zeng W;
XX
DR
    WPI; 2002-674803/72.
XX
    Diagnosing breast cancer in a patient comprises detecting the level of
PT
PT
    gene expression in cell or tissue samples, where a differential gene
PT
    expression is indicative of breast cancer
XX
    Claim 1; SEQ ID NO 1225; 260pp + Sequence Listing; English.
PS
XX
CC
    The present invention relates to methods of diagnosing breast cancer in a
CC
    patient, which comprise detecting the level of expression in a tissue
CC
    sample of two or more genes selected from those shown in ABT09867-
    ABT11112, where a differential expression of the genes indicates breast
CC
CC
    cancer. The methods are useful in diagnosing, treating, detecting the
CC
    progression, and in monitoring treatment of breast cancer in patients.
```

```
the onset or progression of breast cancer. The breast cancer genes may be
CC
   used as diagnostic markers for the prediction or identification of the
CC
   malignant state of breast tissue, for confirming the type and progression
CC
   of cancer, and for drug screening and assays. The present sequence is a
CC
CC
   coding sequence of the invention.
CC
   Note: The sequence data for this patent did not form part of the printed
CC
    specification, but was obtained in electronic format directly from WIPO
CC
   at ftp.wipo.int/pub.published pct sequences.
XX
   Sequence 7260 BP; 2330 A; 1415 C; 1240 G; 2275 T; 0 other;
SO
 Query Match
                    63.9%; Score 334.4; DB 24; Length 7260;
 Best Local Similarity 84.6%; Pred. No. 5.7e-88;
 Matches 445; Conservative 0; Mismatches 26; Indels
                                                 55; Gaps
                                                           4;
         1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 60
Qу
           Db
       311 GGACCGGAGACGCTCTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGGAGAC 370
        61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
Qу
           371 AGGGGCTTTTATTTCAACAAGCCCACAGGGTATGGCTCCAGCAGTCGGAGGGCGCCTCAG 430
Db
       121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Qу
           431 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTAAGGAGGCTGGAGATGTAT 490
Db
       Qу
           491 TGCGCACCCCTCAAGCCTGCCAAGTCAGCTCGCTCTGTCCGTGCCCAGCGCCACACCGAC 550
Db
       241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
Qу
           551 ATGCCCAAGACCCAG----- 565
Dh
       301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
Qу
                566 -----AAGGAAGTACATTTGAAGAACGCAAGTAGAGGGAGTGCAGGAAACAAGAACTA 618
Dh
       361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Qy
           619 CAGGATGTAGGAAGACCCTCCTGAGGAGTGAAGAGTGACATGCCACCGCAGGATCCTTTG 678
Db
       421 CTCTGCAC-AGTTACCTG-TAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACAT 478
Qу
           679 CTCTGCACGAGTTACCTGTTAAACTTTGGAACACCTACCAAAAAATAAGTTTGATAACAT 738
Db
       479 TTCAAAGAT-GGCATTTCCCCCCAATGAAATACACAAGTAAACATTC 523
Qу
           Db
       739 TTAAAAGATGGGCGTTTCCCCCCAATGAAATACACAAGTAAACATTC 784
```

The methods are also useful as a screening tool for agents that modulate

ABK84583 standard; cDNA; 7260 BP.

CC

AC ABK84583; XX DT 14-AUG-2002 (first entry) XX Human cDNA differentially expressed in granulocytic cells #1154. DE XXHuman; ss; granulocytic cell; DNA chip; bacterial infection; KW viral infection; parasitic infection; protozoal infection; KW fungal infection; sterile inflammatory disease; psoriasis; KW KW rheumatoid arthritis; glomerulonephritis; asthma; thrombosis; KW cardiac reperfusion injury; renal reperfusion injury; ARDS; KW adult respiratory distress syndrome; inflammatory bowel disease; KW Crohn's disease; ulcerative colitis; periodontal disease; granulocyte activation; chronic inflammation; allergy. KW XX OS Homo sapiens. XX PNWO200228999-A2. XX PD 11-APR-2002. XX PF03-OCT-2001; 2001WO-US30821. XX PR03-OCT-2000; 2000US-237189P. XX PΑ (GENE-) GENE LOGIC INC. XXPIBeazer-Barclay Y, Weissman SM, Yamaga S, Vockley J; XX WPI; 2002-435328/46. DR XX PTDetecting granulocyte activation by detecting differential expression РΤ of genes associated with granulocyte activation, which serves as PTdiagnostic markers that is useful for monitoring disease states and PTdrug toxicity XXPS Claim 1; SEQ ID No 1154; 114pp; English. XXCCThe invention relates to detecting (M1) granulocyte (GC) activation CC (GCA), by detecting the level of expression of gene(s) (Gs) identified by CC DNA chip analysis as given in the specification, and comparing CC the expression level to an expression level in an unactivated GC, where differential expression of Gs is indicative of GCA. CC CC Also included are modulating (M2) GA by contacting GC with an agent CCthat alters the expression of at least one gene in Gs; (2) screening (M3) CCfor an agent capable of modulating GCA or an inflammation (especially CC chronic) in a tissue, an allergic response in a subject, exposure of a CC subject to a pathogen or sterile inflammatory disease using the CC gene expression profile; (3) detecting (M4) an inflammation (especially CC chronic) in a tissue, an allergic response in a subject, exposure of a CC subject to a pathogen or sterile inflammatory disease, by detecting the CC level of expression in a sample of the tissue of gene(s) from Gs, where CC the level of expression of the gene is indicative of inflammation; CC (4) treating (M5) an inflammation (especially chronic) or in a tissue, CC an allergic response in a subject, exposure of a subject to a pathogen CC or sterile inflammatory disease, by contacting a tissue having

inflammation with an agent that modulates the expression of gene(s)

CC

```
from Gs in the tissue. M1 is useful for detecting GCA; M2 is useful for
CC
CC
    modulating GA; M3 is useful for screening an agent capable of modulating
CC
    GCA preferably in an inflammation in a tissue; M4 is useful for
CC
    detecting an inflammation (especially chronic) in a tissue, an allergic
CC
    response in a subject, exposure of a subject to a pathogen or sterile
CC
    inflammatory disease (e.g. psoriasis, rheumatoid arthritis,
    glomerulonephritis, asthma, thrombosis, cardiac reperfusion injury, renal
CC
CC
    reperfusion injury, ARDS, adult respiratory distress syndrome,
CC
    inflammatory bowel disease, Crohn's disease, ulcerative colitis,
    periodontal disease; also bacterial infection, viral infection,
CC
CC
    parasitic infection, protozoal infection, fungal infection and M5 is
    useful for treating one of the above conditions. The present
CC
    sequence represents a gene differentially expressed in granulocytes.
CC
CC
    Note: The sequence data for this patent did not form part
CC
    of the printed specification, but was obtained in electronic
CC
    format directly from WIPO at
CC
    ftp.wipo.int/pub/published pct sequences.
XX
    Sequence 7260 BP; 2330 A; 1415 C; 1240 G; 2275 T; 0 other;
SO
                     63.9%; Score 334.4; DB 24; Length 7260;
 Query Match
 Best Local Similarity 84.6%; Pred. No. 5.7e-88;
 Matches 445; Conservative
                         0; Mismatches
                                         26; Indels
                                                                4;
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Ov
            Db
        311 GGACCGGAGACGCTCTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 370
         61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
QУ
            371 AGGGGCTTTTATTTCAACAAGCCCACAGGGTATGGCTCCAGCAGTCGGAGGGCGCCTCAG 430
Db
Qу
        121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
            431 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTAAGGAGGCTGGAGATGTAT 490
Db
        Qу
            491 TGCGCACCCTCAAGCCTGCCAAGTCAGCTCGCTCTGTCCGTGCCCAGCGCCACACCGAC 550
Db
        241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
Qу
            551 ATGCCCAAGACCCAG---
Db
        301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
Qу
                 566 -----AAGGAAGTACATTTGAAGAACGCAAGTAGAGGGAGTGCAGGAAACAAGAACTA 618
Db
        361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Qу
            619 CAGGATGTAGGAAGACCCTCCTGAGGAGTGAAGAGTGACATGCCACCGCAGGATCCTTTG 678
Db
        421 CTCTGCAC-AGTTACCTG-TAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACAT 478
Qу
            679 CTCTGCACGAGTTACCTGTTAAACTTTGGAACACCTACCAAAAAATAAGTTTGATAACAT 738
Db
        479 TTCAAAGAT-GGCATTTCCCCCCAATGAAATACACAAGTAAACATTC 523
Qу
```



```
RESULT 14
ABN97244
ΙD
     ABN97244 standard; DNA; 7260 BP.
AC.
     ABN97244;
XX
DT
     13-AUG-2002 (first entry)
XX
DE
     Gene #3742 used to diagnose liver cancer.
XX
KW
     Gene; liver cancer; ds; hepatocellular carcinoma; hepatotropic;
KW
     metastatic liver tumour; cytostatic; expression profile; disease state;
KW
     disease progression; drug toxicity; drug efficacy; drug metabolism.
XX
OS
     Homo sapiens.
ХX
PN
     WO200229103-A2.
XX
PD
     11-APR-2002.
XX
PF
     02-OCT-2001; 2001WO-US30589.
XX
PR
     02-OCT-2000; 2000US-237054P.
XX
PΑ
     (GENE-) GENE LOGIC INC.
XX
PΤ
     Horne D, Alvares C, Peres-Da-Silva S, Vockley JG;
XX
DR
     WPI; 2002-426119/45.
XX
РΤ
     Diagnosing and detecting the progression of liver cancer,
PT
     hepatocellular carcinoma or metastatic liver tumor in a patient,
РΤ
     involves detecting the level of expression of two or more genes in a
PТ
     liver tissue sample
XX
PS
     Claim 1; SEQ ID NO 3742; 298pp; English.
XX
CC
     The invention relates to a novel method for diagnosing and detecting the
CC
     progression of liver cancer, hepatocellular carcinoma or metastatic liver
CC
     tumour in a patient, and differentiating metastatic liver cancer from
CC
     hepatocellular carcinoma in a patient, involving detecting the level of
CC
     expression of two or more genes represented in ABN93503-ABN97455 in a
CC
     tissue sample. The method of the invention has hepatotropic, and
CC
     cytostatic activity. The method is useful for diagnosing and detecting
CC
     the progression of liver cancer, hepatocellular carcinoma and metastatic
CC
     liver carcinoma in a patient. The method is useful for identifying
CC
     expression profiles which serve as useful diagnostic markers as well as
CC
     markers that can be used to monitor disease states, disease progression,
CC
     drug toxicity, drug efficacy and drug metabolism.
     Note: The sequence data for this patent did not form part of the printed
CC
CC
     specification, but was obtained in electronic format directly from WIPO
CC
     at ftp.wipo.int/pub/published pct sequences.
XX
```

```
Sequence 7260 BP; 2330 A; 1415 C; 1240 G; 2275 T; 0 other;
SO
                   63.9%; Score 334.4; DB 24; Length 7260;
                   84.6%; Pred. No. 5.7e-88;
 Best Local Similarity
 Matches 445; Conservative
                        0; Mismatches
                                     26; Indels
         1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 60
Qу
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       311 GGACCGGAGACGCTCTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGGAGAC 370
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QУ
           Db
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       121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
QУ
           Db
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Qу
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Db
       361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
QУ
           619 CAGGATGTAGGAAGACCCTCCTGAGGAGTGAAGAGTGACATGCCACCGCAGGATCCTTTG 678
Db
       421 CTCTGCAC-AGTTACCTG-TAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACAT 478
Qу
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Db
       479 TTCAAAGAT-GGCATTTCCCCCAATGAAATACACAAGTAAACATTC 523
Qу
       Db
RESULT 15
ABK64812
    ABK64812 standard; DNA; 7260 BP.
ID
XX
AC
   ABK64812;
XX
DT
    18-JÚN-2002 (first entry)
XX
```

```
Human benign prostatic hyperplasia gene #707.
DΕ
XX
KW
     Human; benign prostatic hyperplasia; BPH; prostate cancer; gene; ds.
XX
OS
     Homo sapiens.
XX
```

```
PN
    WO200212440-A2.
XX
PD
    14-FEB-2002.
XX
ΡF
    07-AUG-2001; 2001WO-US24708.
XX
PR
     07-AUG-2000; 2000US-223323P.
PR
     05-JUN-2001; 2001US-0873319.
XX
PΑ
     (GENE-) GENE LOGIC INC.
     (NISB ) JAPAN TOBACCO INC.
PA
XX
    Munger WE, Kulkarni P, Getzenberg RH, Waga I, Yamamoto J;
PΙ
XX
DR
    WPI; 2002-257476/30.
XX
РТ
     Identifying drugs for and diagnosing benign prostatic hyperplasia, by
     detecting expression levels of one or more genes in prostate cells from
PT
PT
     patient that are differentially regulated compared to normal prostate
PT
     cells -
XX
PS
     Disclosure; Page 391-393; 444pp; English.
XX
CC
     The invention relates to a method of diagnosing (I) the onset or
CC
     progression of benign prostatic hyperplasia (BPH), or screening (II) for
     or identifying an agent that modulates the onset or progression of BPH.
CC
     The method is based on changes in gene expression in BPH tissue isolated
CC
CC
     from patients exhibiting different clinical states of prostate
     hyperplasia as compared to normal prostate tissue. (I) comprises
CC
CC
     detecting the expression levels of one or more genes in prostate cells
CC
     from the subject that are differentially regulated compared to normal
CC
     prostate cells. (II) comprises preparing a first gene expression profile
CC
     of BPH cells or BPH-like cell population, exposing the cells to the
     agent, preparing a second gene expression profile of the agent exposed
CC
CC
     cells, and comparing the first and second gene expression profiles.
     (I) is useful for diagnosing the onset or progression of BPH. (II) is
CC
CC
     useful for identifying an agent that modulates the onset or progression
CC
     of BPH. The methods are useful to present information identifying
CC
     the expression level in a tissue or cells, by comparing the expression
     level of genes given in the specification in the tissue or cells to the
CC
CC
     level of expression of gene in the database, and displaying the
CC
     expression levels of at least one gene in the tissue or cell sample
     compared to the expression level in BPH. Agents using (II) are useful for
CC
CC
     treating BPH or prostate cancer. ABK64106-ABK64860 represent human
     benign prostatic hyperplasia gene sequences of the invention.
CC
XX
     Sequence 7260 BP; 2330 A; 1415 C; 1240 G; 2275 T; 0 other;
SO
                          63.9%; Score 334.4; DB 24; Length 7260;
  Query Match
                          84.6%;
                                 Pred. No. 5.7e-88;
  Best Local Similarity
                                0; Mismatches
                                                 26; Indels
                                                               55; Gaps
                                                                            4;
  Matches 445; Conservative
            {\tt 1} \quad {\tt GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGAGAC} \quad {\tt 60} \\
Qу
              Db
          311 GGACCGGAGACGCTCTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 370
           61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
Qу
```

Db	371	AGGGGCTTTTATTTCAACAAGCCCACAGGGTATGGCTCCAGCAGTCGGAGGGCGCCTCAG	430
Qу	121	ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC	180
Db	431	ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTAAGGAGGCTGGAGATGTAT	490
Qу	181	TGTGCACCCCTCAAGCCGGCAAAGGCAGCCCGCTCCGTCCG	240
Db	491	TGCGCACCCCTCAAGCCTGCCAAGTCAGCTCGCTCTGTCCGTGCCCAGCGCCACACCGAC	550
Qу	241	ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG	300
Db	551	ATGCCCAAGACCCAG	565
Qу	3'01	AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA	360
Db	566		618
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Db	619	CAGGATGTAGGAAGACCCTCCTGAGGAGTGAAGAGTGACATGCCACCGCAGGATCCTTTG	678
Qy	421	CTCTGCAC-AGTTACCTG-TAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACAT	478
Db	679	CTCTGCACGAGTTACCTGTTAAACTTTGGAACACCTACCAAAAAATAAGTTTGATAACAT	738
Qу	479	TTCAAAGAT-GGCATTTCCCCCAATGAAATACACAAGTAAACATTC 523	
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OM nucleic - nucleic search, using sw model

Run on:

December 13, 2003, 06:03:55; Search time 48.3585 Seconds

(without alignments)

4773.589 Million cell updates/sec

Title:

US-09-852-261-5

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Scoring table:

IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched:

569978 seqs, 220691566 residues

Total number of hits satisfying chosen parameters:

1139956

Minimum DB seq length: 0

Maximum DB seg length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

Issued Patents NA:*

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- 3: /cgn2_6/ptodata/1/ina/6A_COMB.seq:*
- 4: /cgn2 6/ptodata/1/ina/6B_COMB.seq:*
- /cgn2_6/ptodata/1/ina/PCTUS_COMB.seq:*
- /cgn2 6/ptodata/1/ina/backfiles1.seq:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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1	523	100.0	553	3	US-09-142-583A-3	Sequence 3, Appli
2	- 523	100.0	553	3	US-09-142-583A-5	Sequence 5, Appli
. 3	332.8	63.6	777	3	US-09-142-583A-10	Sequence 10, Appl
4	331.2	63.3	622	6	5405942-2	Patent No. 5405942
5	274.6	52.5	5707	2	US-08-472-809B-8	Sequence 8, Appli
.6	274.6	52.5	6345	2	US-08-472-809B-7	Sequence 7, Appli
7	234.4	44.8	357	6	5405942-13	Patent No. 5405942
8	232.8	44.5	357	. 6	5405942-9	Patent No. 5405942.
9	191.4	36.6	210	6	5405942-7	Patent No. 5405942
10	191.4	36.6	210	6	5405942-11	Patent No. 5405942
11	191.4	36.6	2862	4	US-09-255-829-13	Sequence 13, Appl

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             35.4
                      390 3
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ALIGNMENTS

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RESULT 1
US-09-142-583A-3
; Sequence 3, Application US/09142583A
 Patent No. 6221842
   GENERAL INFORMATION:
         APPLICANT: GOLDSPINK, GEOFFREY
         TITLE OF INVENTION: METHOD OF TREATING MUSCULAR DISORDERS
         NUMBER OF SEQUENCES: 11
         CORRESPONDENCE ADDRESS:
              ADDRESSEE: NIXON & VANDERHYE P.C.
              STREET: 1100 NORTH GLEBE ROAD
              CITY: ARLINGTON
              STATE: VA
              COUNTRY: USA
              ZIP: 22201
         COMPUTER READABLE FORM:
              MEDIUM TYPE: Floppy disk
              COMPUTER: IBM PC compatible
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OPERATING SYSTEM: PC-DOS/MS-DOS
          SOFTWARE: PatentIn Release #1.0, Version #1.25
      CURRENT APPLICATION DATA:
          APPLICATION NUMBER: US/09/142,583A
          FILING DATE: 29-Oct-1998
          CLASSIFICATION: <Unknown>
      PRIOR APPLICATION DATA:
          APPLICATION NUMBER: WO PCT/GB97/00658
          FILING DATE: 11-MAR-1997
          APPLICATION NUMBER: GB 9605124.8
          FILING DATE: 11-MAR-1996
      ATTORNEY/AGENT INFORMATION:
          NAME: SADOFF, B. J.
          REGISTRATION NUMBER: 36663
          REFERENCE/DOCKET NUMBER: 117-263
      TELECOMMUNICATION INFORMATION:
          TELEPHONE: 7038164000
          TELEFAX: 7038164100
   INFORMATION FOR SEQ ID NO: 3:
      SEQUENCE CHARACTERISTICS:
          LENGTH: 553 base pairs
          TYPE: nucleic acid
          STRANDEDNESS: both
          TOPOLOGY: linear
      MOLECULE TYPE: cDNA
      FEATURE:
          NAME/KEY: CDS
          LOCATION: 1..363
      SEQUENCE DESCRIPTION: SEQ ID NO: 3:
US-09-142-583A-3
 Query Match
                    100.0%; Score 523; DB 3; Length 553;
                   100.0%; Pred. No. 1.6e-154;
 Best Local Similarity
                        0; Mismatches
 Matches 523; Conservative
                                      0; Indels
                                                    Gaps
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             451 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTT 510
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Qу
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RESULT 2
US-09-142-583A-5
; Sequence 5, Application US/09142583A
; Patent No. 6221842
   GENERAL INFORMATION:
        APPLICANT: GOLDSPINK, GEOFFREY
        TITLE OF INVENTION: METHOD OF TREATING MUSCULAR DISORDERS
        NUMBER OF SEQUENCES: 11
        CORRESPONDENCE ADDRESS:
             ADDRESSEE: NIXON & VANDERHYE P.C.
             STREET: 1100 NORTH GLEBE ROAD
             CITY: ARLINGTON
             STATE: VA
             COUNTRY: USA
             ZIP: 22201
        COMPUTER READABLE FORM:
             MEDIUM TYPE: Floppy disk
             COMPUTER: IBM PC compatible
             OPERATING SYSTEM: PC-DOS/MS-DOS
             SOFTWARE: PatentIn Release #1.0, Version #1.25
        CURRENT APPLICATION DATA:
             APPLICATION NUMBER: US/09/142,583A
             FILING DATE: 29-Oct-1998
             CLASSIFICATION: <Unknown>
        PRIOR APPLICATION DATA:
             APPLICATION NUMBER: WO PCT/GB97/00658
             FILING DATE: 11-MAR-1997
             APPLICATION NUMBER: GB 9605124.8
             FILING DATE: 11-MAR-1996
        ATTORNEY/AGENT INFORMATION:
             NAME: SADOFF, B. J.
             REGISTRATION NUMBER: 36663
             REFERENCE/DOCKET NUMBER: 117-263
        TELECOMMUNICATION INFORMATION:
             TELEPHONE: 7038164000
             TELEFAX: 7038164100
    INFORMATION FOR SEQ ID NO: 5:
        SEQUENCE CHARACTERISTICS:
             LENGTH: 553 base pairs
             TYPE: nucleic acid
             STRANDEDNESS: both
             TOPOLOGY: linear
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MOLECULE TYPE: cDNA
      FEATURE:
          NAME/KEY: CDS
          LOCATION: 341..397
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US-09-142-583A-5
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 Best Local Similarity
                  100.0%; Pred. No. 1.6e-154;
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 Matches 523; Conservative
                                    0; Indels
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RESULT 3
US-09-142-583A-10
 Sequence 10, Application US/09142583A
 Patent No. 6221842
   GENERAL INFORMATION:
      APPLICANT: GOLDSPINK, GEOFFREY
      TITLE OF INVENTION: METHOD OF TREATING MUSCULAR DISORDERS
      NUMBER OF SEQUENCES: 11
      CORRESPONDENCE ADDRESS:
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ADDRESSEE: NIXON & VANDERHYE P.C.
            STREET: 1100 NORTH GLEBE ROAD
            CITY: ARLINGTON
            STATE: VA
            COUNTRY: USA
            ZIP: 22201
       COMPUTER READABLE FORM:
            MEDIUM TYPE: Floppy disk
            COMPUTER: IBM PC compatible
            OPERATING SYSTEM: PC-DOS/MS-DOS
            SOFTWARE: PatentIn Release #1.0, Version #1.25
       CURRENT APPLICATION DATA:
            APPLICATION NUMBER: US/09/142,583A
            FILING DATE: 29-Oct-1998
            CLASSIFICATION: <Unknown>
       PRIOR APPLICATION DATA:
            APPLICATION NUMBER: WO PCT/GB97/00658
            FILING DATE: 11-MAR-1997
            APPLICATION NUMBER: GB 9605124.8
            FILING DATE: 11-MAR-1996
       ATTORNEY/AGENT INFORMATION:
            NAME: SADOFF, B. J.
            REGISTRATION NUMBER: 36663
            REFERENCE/DOCKET NUMBER: 117-263
       TELECOMMUNICATION INFORMATION:
            TELEPHONE: 7038164000
            TELEFAX: 7038164100
   INFORMATION FOR SEQ ID NO: 10:
       SEQUENCE CHARACTERISTICS:
            LENGTH: 777 base pairs
            TYPE: nucleic acid
            STRANDEDNESS: both
            TOPOLOGY: linear
       MOLECULE TYPE: cDNA
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            NAME/KEY: CDS
            LOCATION: 26..493
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 Best Local Similarity 84.4%; Pred. No. 9.8e-95;
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RESULT 4
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; Patent No. 5405942
    APPLICANT: BELL, GRAEME I.; RALL, LESLIE B.; MERRYWEATHER,
; JAMES P.
    TITLE OF INVENTION: PREPRO INSULIN-LIKE GROWTH FACTORS
; I AND II
    NUMBER OF SEQUENCES: 16
    CURRENT APPLICATION DATA:
     APPLICATION NUMBER: US/07/65,673
     FILING DATE: 16-JUN-1987
    PRIOR APPLICATION DATA:
     APPLICATION NUMBER: 630,557
     FILING DATE: 19-JUL-1984
;SEQ ID NO:2:
     LENGTH: 622
5405942-2
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                    63.3%; Score 331.2; DB 6; Length 622;
                    68.3%; Pred. No. 2.8e-94;
 Best Local Similarity
 Matches 359; Conservative 84; Mismatches 28;
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        121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
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       421 CTCTGCAC-AGTTACCTG-TAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACAT 478
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           Db
       479 TTCAAAGAT-GGCATTTCCCCCAATGAAATACACAAGTAAACATTC 523
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           473 UUAAAAGAUGGGCGUUUCCCCCAAUGAAAUACACAAGUAAACAUUC 518
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RESULT 5
US-08-472-809B-8
; Sequence 8, Application US/08472809B
 Patent No. 5925564
  GENERAL INFORMATION:
    APPLICANT: Schwartz, Robert J.
    APPLICANT: DeMayo, Franco J.
    APPLICANT: O'Malley, Bert W.
    TITLE OF INVENTION: Expression Vector Systems and
    TITLE OF INVENTION: Method of Use
    NUMBER OF SEQUENCES: 8
    CORRESPONDENCE ADDRESS:
     ADDRESSEE: Lyon & Lyon
     STREET: 633 West Fifth Street
     STREET: Suite 4700
     CITY: Los Angeles
     STATE: California
     COUNTRY: U.S.A.
     ZIP: 90071-2066
    COMPUTER READABLE FORM:
     MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
     MEDIUM TYPE: storage
     COMPUTER: IBM Compatible
     OPERATING SYSTEM: IBM P.C. DOS 5.0
     SOFTWARE: Word Perfect 5.1
    CURRENT APPLICATION DATA:
     APPLICATION NUMBER: US/08/472,809B
     FILING DATE: June 7, 1995
     CLASSIFICATION: 435
    PRIOR APPLICATION DATA:
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APPLICATION NUMBER: 08/209,846

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FILING DATE: March 9, 1994
     APPLICATION NUMBER: 07/789,919
     FILING DATE: No. 5925564ember 6, 1991
   ATTORNEY/AGENT INFORMATION:
     NAME: Warburg, Richard J.
     REGISTRATION NUMBER: 32,327
     REFERENCE/DOCKET NUMBER: 214/212
   TELECOMMUNICATION INFORMATION:
     TELEPHONE: (213) 489-1600
     TELEFAX: (213) 955-0440
     TELEX: 67-3510
  INFORMATION FOR SEO ID NO:
   SEQUENCE CHARACTERISTICS:
     LENGTH: 5707 bases
     TYPE: nucleic acid
     STRANDEDNESS: double
     TOPOLOGY: linear
   MOLECULE TYPE: cDNA
US-08-472-809B-8
                    52.5%; Score 274.6; DB 2; Length 5707;
 Query Match
 Best Local Similarity 82.2%; Pred. No. 5.3e-76;
 Matches 351; Conservative 0; Mismatches
                                           Indels
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           793 GGACCGGAGACGCTCTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGGAGAC 852
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            11111
       1161 GGCTGCA 1167
Db
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RESULT 6
US-08-472-809B-7
; Sequence 7, Application US/08472809B
; Patent No. 5925564
  GENERAL INFORMATION:
    APPLICANT: Schwartz, Robert J.
    APPLICANT: DeMayo, Franco J.
    APPLICANT: O'Malley, Bert W.
    TITLE OF INVENTION: Expression Vector Systems and
    TITLE OF INVENTION: Method of Use
    NUMBER OF SEQUENCES: 8
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: Lyon & Lyon
      STREET: 633 West Fifth Street
      STREET: Suite 4700
      CITY: Los Angeles
      STATE: California
      COUNTRY: U.S.A.
      ZIP: 90071-2066
    COMPUTER READABLE FORM:
      MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
      MEDIUM TYPE: storage
      COMPUTER: IBM Compatible
      OPERATING SYSTEM: IBM P.C. DOS 5.0
      SOFTWARE: Word Perfect 5.1
    CURRENT APPLICATION DATA:
      APPLICATION NUMBER: US/08/472,809B
      FILING DATE: June 7, 1995
      CLASSIFICATION: 435
    PRIOR APPLICATION DATA:
      APPLICATION NUMBER: 08/209,846
      FILING DATE: March 9, 1994
      APPLICATION NUMBER: 07/789,919
      FILING DATE: No. 5925564ember 6, 1991
    ATTORNEY/AGENT INFORMATION:
      NAME: Warburg, Richard J.
      REGISTRATION NUMBER: 32,327
      REFERENCE/DOCKET NUMBER: 214/212
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: (213) 489-1600
      TELEFAX: (213) 955-0440
      TELEX: 67-3510
   INFORMATION FOR SEQ ID NO: 7:
    SEQUENCE CHARACTERISTICS:
      LENGTH: 6345 bases
      TYPE: nucleic acid
      STRANDEDNESS: double
      TOPOLOGY: linear
    MOLECULE TYPE: cDNA
US-08-472-809B-7
 Query Match
                        52.5%; Score 274.6; DB 2;
                                                   Length 6345;
 Best Local Similarity 82.2%; Pred. No. 5.6e-76;
 Matches 351; Conservative 0; Mismatches 24; Indels
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           Db
       3762 AGGGGCTTTTATTTCAACAAGCCCACAGGGTATGGCTCCAGCAGTCGGAGGGCGCCTCAG 3821
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        QУ
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   APPLICANT: BELL, GRAEME I.; RALL, LESLIE B.; MERRYWEATHER.
    TITLE OF INVENTION: PREPRO INSULIN-LIKE GROWTH FACTORS
; I AND II
   NUMBER OF SEQUENCES: 16
    CURRENT APPLICATION DATA:
     APPLICATION NUMBER: US/07/65,673
     FILING DATE: 16-JUN-1987
    PRIOR APPLICATION DATA:
     APPLICATION NUMBER: 630,557
     FILING DATE: 19-JUL-1984
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; Patent No. 5405942
   APPLICANT: BELL, GRAEME I.; RALL, LESLIE B.; MERRYWEATHER,
   TITLE OF INVENTION: PREPRO INSULIN-LIKE GROWTH FACTORS
; I AND II
   NUMBER OF SEQUENCES: 16
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     APPLICATION NUMBER: US/07/65,673
     FILING DATE: 16-JUN-1987
    PRIOR APPLICATION DATA:
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     FILING DATE: 19-JUL-1984
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; Patent No. 5405942
    APPLICANT: BELL, GRAEME I.; RALL, LESLIE B.; MERRYWEATHER.
    TITLE OF INVENTION: PREPRO INSULIN-LIKE GROWTH FACTORS
; I AND II
    NUMBER OF SEQUENCES: 16
    CURRENT APPLICATION DATA:
      APPLICATION NUMBER: US/07/65,673
      FILING DATE: 16-JUN-1987
    PRIOR APPLICATION DATA:
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      FILING DATE: 19-JUL-1984
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; Patent No. 5405942
    APPLICANT: BELL, GRAEME I.; RALL, LESLIE B.; MERRYWEATHER,
    TITLE OF INVENTION: PREPRO INSULIN-LIKE GROWTH FACTORS
; I AND II
    NUMBER OF SEQUENCES: 16
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      APPLICATION NUMBER: US/07/65,673
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; Sequence 13, Application US/09255829
; Patent No. 6461617
  GENERAL INFORMATION:
    APPLICANT: Shone, Clifford Charles
    APPLICANT: Quinn, Conrad Padraig
    APPLICANT: Foster, Keith Alan
    TITLE OF INVENTION: Recombinant Toxin Fragments
    NUMBER OF SEQUENCES: 29
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: STERNE, KESSLER, GOLDSTEIN, & FOX P.L.L.C.
      STREET: 1100 NEW YORK AVENUE, NW, SUITE 600
      CITY: WASHINGTON
      STATE: DC
     COUNTRY: USA
      ZIP: 20005-3934
    COMPUTER READABLE FORM:
      MEDIUM TYPE: Floppy disk
      COMPUTER: IBM PC compatible
      OPERATING SYSTEM: PC-DOS/MS-DOS
      SOFTWARE: PatentIn Release #1.0, Version #1.30 (EPO)
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      FILING DATE: 23-FEB-1999
    PRIOR APPLICATION DATA:
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      FILING DATE: 22-AUG-1997
    PRIOR APPLICATION DATA:
      APPLICATION NUMBER: US 08/782,893
      FILING DATE: 27-DEC-1996
    ATTORNEY/AGENT INFORMATION:
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LENGTH: 210

NAME: ESMOND, ROBERT W.

5405942-11

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REGISTRATION NUMBER: 32,893
     REFERENCE/DOCKET NUMBER: 1581.0130002
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: 202-371-2600
      TELEFAX: 202-371-2540
  INFORMATION FOR SEQ ID NO: 13:
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; Patent No. 5405942
    APPLICANT: BELL, GRAEME I.; RALL, LESLIE B.; MERRYWEATHER,
    TITLE OF INVENTION: PREPRO INSULIN-LIKE GROWTH FACTORS
; I AND II
    NUMBER OF SEQUENCES: 16
    CURRENT APPLICATION DATA:
      APPLICATION NUMBER: US/07/65,673
      FILING DATE: 16-JUN-1987
    PRIOR APPLICATION DATA:
      APPLICATION NUMBER: 630,557
     FILING DATE: 19-JUL-1984
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; Sequence 1, Application US/08308196A
; Patent No. 5612198
  GENERAL INFORMATION:
  APPLICANT: Brierley, Russell A. APPLICANT: Davis, Geneva R.
    APPLICANT: Holtz, Gregory C.
    APPLICANT: Gleeson, Martin A.
    APPLICANT: Howard, Bradley D.
    TITLE OF INVENTION: Production of Insulin-Like Growth
    TITLE OF INVENTION: Factor-1 in Methylotrophic Yeast Cells
    NUMBER OF SEQUENCES: 17
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: Brown, Martin, Haller & McClain
      STREET: 1660 Union Street
      CITY: San Diego
      STATE: California
      COUNTRY: USA
      ZIP: 92101-2926
    COMPUTER READABLE FORM:
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      COMPUTER: IBM PC compatible
      OPERATING SYSTEM: PC-DOS/MS-DOS
      SOFTWARE: PatentIn Release #1.0, Version #1.25
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      FILING DATE: 09-SEPT-1994
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      APPLICATION NUMBER: US 07/983,523
      FILING DATE: 03-MAR-1993
    PRIOR APPLICATION DATA:
      APPLICATION NUMBER: US 07/578,728
      FILING DATE: 04-SEP-1990
    ATTORNEY/AGENT INFORMATION:
      NAME: Seidman, Stephanie L.
      REGISTRATION NUMBER: 33,779
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TELECOMMUNICATION INFORMATION:
      TELEPHONE: (619)238-0999
      TELEFAX: (619)238-0062
  INFORMATION FOR SEQ ID NO:
    SEQUENCE CHARACTERISTICS:
      LENGTH: 240 base pairs
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  GENERAL INFORMATION:
    APPLICANT: Brierley, Russell A.
    APPLICANT: Davis, Geneva R.
    APPLICANT: Holtz, Gregory C.
    APPLICANT: Gleeson, Martin A.
    APPLICANT: Bradley, D. H.
    TITLE OF INVENTION: Production of Insulin-Like Growth
TITLE OF INVENTION: Factor-1 in Methylotrophic Yeast Cells
    NUMBER OF SEQUENCES: 12
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: Fitch, Even, Tabin & Flannery
      STREET: 135 South LaSalle Street, Suite 900
      CITY: Chicago
      STATE: Illinois
      COUNTRY: USA
      ZIP: 60603
    COMPUTER READABLE FORM:
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MEDIUM TYPE: Floppy disk
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      FILING DATE: 19910409
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      FILING DATE: 04-SEP-1990
    ATTORNEY/AGENT INFORMATION:
      NAME: Seidman, Stephanie L.
      REGISTRATION NUMBER: 33,779
      REFERENCE/DOCKET NUMBER: 51874
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: (619) 552-1311
      TELEFAX: (619)552-0095
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; Patent No. 6107057
  GENERAL INFORMATION:
    APPLICANT: Crawford, Kenneth
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APPLICANT: Zaror, Isabel
    APPLICANT: Innis, Michael
    TITLE OF INVENTION: Pichia Secretory Leader for Protein
    TITLE OF INVENTION: Expression
    NUMBER OF SEQUENCES: 40
    CORRESPONDENCE ADDRESS:
     ADDRESSEE: Chiron Corporation
     STREET: 4560 Horton Street
     CITY: Emeryville
     STATE: California
     COUNTRY: United States
      ZIP: 94608
    COMPUTER READABLE FORM:
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    ATTORNEY/AGENT INFORMATION:
      NAME: Chung, Ling-Fong
      REGISTRATION NUMBER: 36,482
      REFERENCE/DOCKET NUMBER: 1165.100
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: (510) 601-2704
      TELEFAX: (510) 655-3542
  INFORMATION FOR SEQ ID NO: 13:
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Title:

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	26	127	24.3	621	9	US-09-921-398-40	Sequence 40, Appl
	27	127	24.3	621	15	US-10-280-826-40	Sequence 40, Appl
	28	113.4	21.7	480	9	US-09-921-398-38	Sequence 38, Appl
	29	113.4	21.7	480	15	US-10-280-826-38	Sequence 38, Appl
	30	101.8	19.5	210	13	US-09-807-742-19	Sequence 19, Appl
	31	77.2	14.8	854	10	US-09-954-531-989	Sequence 989, App
C	32	75.4	14.4	447	9	US-09-922-217-917	Sequence 917, App
C	33	75.4	14.4	447	10	US-09-833-263-917	Sequence 917, App
C	34	75.4	14.4	447	14	US-10-025-380-917	Sequence 917, App
C	35	75.2	14.4	437	15	US-10-066-543-663	Sequence 663, App
C	36	75.2	14.4	493	15	US-10-066-543-997	Sequence 997, App
C	37	75.2	14.4	518	15	US-10-066-543-1040	Sequence 1040, Ap
C	38	75.2	14.4	536	15	US-10-066-543 - 428	Sequence 428, App
	39	75.2	14.4	543	15		Sequence 1, Appli
С	40	75.2	14.4	549	15	US-10-066-543-478	Sequence 478, App
C	41	75.2	14.4	574	9	US-09-922-217-918	Sequence 918, App
C	42	75.2	14.4	574	10	US-09-833-263-918	Sequence 918, App
C	43	75.2	14.4	574	14	US-10-025-380-918	Sequence 918, App
C	44	75.2	14.4	577	15	US-10-066-543 - 1137	Sequence 1137, Ap
С	45	75.2	14.4	579	15	US-10-066-543-1094	Sequence 1094, Ap

ALIGNMENTS

RESULT 1

US-09-852-261-5

[;] Sequence 5, Application US/09852261 ; Patent No. US20020083477A1

[;] GENERAL INFORMATION:

[;] APPLICANT: GOLDSPINK, GEOFFREY

```
APPLICANT: TERENGHI, GIORGIO
  TITLE OF INVENTION: REPAIR OF NERVE DAMAGE
  FILE REFERENCE: 117-351
  CURRENT APPLICATION NUMBER: US/09/852,261
  CURRENT FILING DATE: 2001-05-10
  PRIOR APPLICATION NUMBER: GB 0011278.9
  PRIOR FILING DATE: 2000-05-10
  NUMBER OF SEQ ID NOS: 14
  SOFTWARE: PatentIn Ver. 2.1
 SEQ ID NO 5
  LENGTH: 523
  TYPE: DNA
  ORGANISM: Oryctolagus cuniculus
US-09-852-261-5
                   100.0%; Score 523; DB 9; Length 523;
 Query Match
 Best Local Similarity
                   100.0%; Pred. No. 1.9e-161;
 Matches 523; Conservative 0; Mismatches
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                                        Indels
                                                   Gaps
                                                         0;
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Qу
          1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGGAGAC 60
Db
        61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
Qу
          61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
Db
       121 ACAGGCATCGTGGATGATGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
QУ
          121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Db
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QУ
          Dh
       241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
Qу
          241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
Db
       301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
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          301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
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       361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Qу
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       421 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTT 480
Qу
          421 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTT 480
Dh
       481 CAAAGATGGCATTTCCCCCAATGAAATACACAAGTAAACATTC 523
Qу
          481 CAAAGATGGCATTTCCCCCCAATGAAATACACAGTAAACATTC 523
Dh
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US-09-852-261-1
; Sequence 1, Application US/09852261
 Patent No. US20020083477A1
 GENERAL INFORMATION:
  APPLICANT: GOLDSPINK, GEOFFREY
  APPLICANT: TERENGHI, GIORGIO
  TITLE OF INVENTION: REPAIR OF NERVE DAMAGE
  FILE REFERENCE: 117-351
  CURRENT APPLICATION NUMBER: US/09/852,261
  CURRENT FILING DATE: 2001-05-10
  PRIOR APPLICATION NUMBER: GB 0011278.9
  PRIOR FILING DATE: 2000-05-10
  NUMBER OF SEQ ID NOS: 14
  SOFTWARE: PatentIn Ver. 2.1
 SEO ID NO 1
   LENGTH: 517
   TYPE: DNA
   ORGANISM: Homo sapiens
US-09-852-261-1
                    89.4%; Score 467.4; DB 9; Length 517;
 Query Match
                    96.2%; Pred. No. 3.8e-143;
 Best Local Similarity
                         0; Mismatches
                                                           2;
                                      16:
                                          Indels
                                                     Gaps
 Matches 501; Conservative
                                                  4:
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           1 GGACCGGAGACGCTCTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 60
Db
        61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
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           61 AGGGGCTTTTATTTCAACAAGCCCACAGGGTATGGCTCCAGCAGTCGGAGGGCGCCTCAG 120
Db
        121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
QУ
           121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTAAGGAGGCTGGAGATGTAT 180
Dh
        Qу
           181 TGCGCACCCTCAAGCCTGCCAAGTCAGCTCGCTCTGTCCGTGCCCAGCGCCACACCGAC 240
Db
        241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
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           241 ATGCCCAAGACCCAGAAGTATCAGCCCCCATCTACCAACAAGAACACGAAGTCTCA---G 297
Db
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Qу
           298 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 357
Db
        361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Qу
           358 CAGGATGTA-GAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 416
Db
        421 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTT 480
Qу
           417 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTT 476
Db
        481 CAAAGATGGCATTTCCCCCCAATGAAATACACAAGTAAACAT 521
Qу
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RESULT 3
US-09-852-261-13
; Sequence 13, Application US/09852261
; Patent No. US20020083477A1
: GENERAL INFORMATION:
  APPLICANT: GOLDSPINK, GEOFFREY
  APPLICANT: TERENGHI, GIORGIO
  TITLE OF INVENTION: REPAIR OF NERVE DAMAGE
  FILE REFERENCE: 117-351
  CURRENT APPLICATION NUMBER: US/09/852,261
  CURRENT FILING DATE: 2001-05-10
  PRIOR APPLICATION NUMBER: GB 0011278.9
  PRIOR FILING DATE: 2000-05-10
  NUMBER OF SEQ ID NOS: 14
  SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 13
   LENGTH: 471
   TYPE: DNA
   ORGANISM: Oryctolagus cuniculus
US-09-852-261-13
                   78.2%; Score 409; DB 9; Length 471;
 Query Match
 Best Local Similarity 90.1%; Pred. No. 6.1e-124;
 Matches 471: Conservative 0; Mismatches 0; Indels
                                               52; Gaps
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          1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 60
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Qу
          61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
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       121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Qу
          121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Db
       Qу
          Db
       241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
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          241 ATGCCCAAGACTCAG-----
Db
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QУ
               256 -----AAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 308
Db
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Qу
          Db.
       309 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 368
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421 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTT 480
Qу
           369 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACATTT 428
Db
       481 CAAAGATGGCATTTCCCCCCAATGAAATACACAAGTAAACATTC 523
QУ
           429 CAAAGATGGCATTTCCCCCCAATGAAATACACAAGTAAACATTC 471
Db
RESULT 4
US-09-852-261-3
; Sequence 3, Application US/09852261
; Patent No. US20020083477A1
; GENERAL INFORMATION:
  APPLICANT: GOLDSPINK, GEOFFREY
  APPLICANT: TERENGHI, GIORGIO
  TITLE OF INVENTION: REPAIR OF NERVE DAMAGE
  FILE REFERENCE: 117-351
  CURRENT APPLICATION NUMBER: US/09/852,261
  CURRENT FILING DATE: 2001-05-10
  PRIOR APPLICATION NUMBER: GB 0011278.9
  PRIOR FILING DATE: 2000-05-10
  NUMBER OF SEQ ID NOS: 14
  SOFTWARE: PatentIn Ver. 2.1
 SEO ID NO 3
   LENGTH: 539
   TYPE: DNA
   ORGANISM: Rattus sp.
US-09-852-261-3
                    68.2%; Score 356.8; DB 9; Length 539;
 Query Match
                    82.3%; Pred. No. 9.9e-107;
 Best Local Similarity
                         0; Mismatches 87;
                                          Indels 7; Gaps
                                                            2;
 Matches 436: Conservative
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           1 GGACCAGAGACCCTTTGCGGGGCTGAGCTGGTGGACGCTCTTCAGTTCGTGTGTGGACCA 60
Dh
         61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
Qу
           61 AGGGGCTTTTACTTCAACAAGCCCACAGTCTATGGCTCCAGCATTCGGAGGGCACCACAG 120
Db
        121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Oy
           121 ACGGCATTGTGGATGAGTGTTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Db
        Qу
                   181 TGTGTCCGCTGCAAGCCTACAAAGTCAGCTCGTTCCATCCGGGCCCAGCGCCCACACTGAC 240
Db
        241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
Qу
           241 ATGCCCAAGACTCAGAAGTCCCAGCCCCTATCGACACACAAGAAAAGGAAGCTGCAAAGG 300
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Db
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                         421 CTGCTTGAGCAACCTGCAAAACATCGGAACACCTGCCAAATATCAATAATGAGTTCAATA 480
Db
        475 ACATTTCAAAGAT-GGCATTTCCCCCAATGAAATACACAAGTAAACATTC 523
QУ
            481 TCATTTCAGAGATGGGCATTTCCCTCAATGAAATACACAAGTAAACATTC 530
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RESULT 5
US-10-161-088-1
; Sequence 1, Application US/10161088
 Publication No. US20030077761A1
; GENERAL INFORMATION:
  APPLICANT: Parrow, Vendela
  APPLICANT: Rosengren, Linda
  TITLE OF INVENTION: NEW METHODS
  FILE REFERENCE: 13425-111001
  CURRENT APPLICATION NUMBER: US/10/161,088
  CURRENT FILING DATE: 2002-05-31
  PRIOR APPLICATION NUMBER: SE 0101934-8
  PRIOR FILING DATE: 2001-06-01
  NUMBER OF SEQ ID NOS: 3
  SOFTWARE: FastSEQ for Windows Version 4.0
 SEO ID NO 1
   LENGTH: 651
   TYPE: DNA
   ORGANISM: Homo sapiens
   FEATURE:
   NAME/KEY: CDS
   LOCATION: (73)...(471)
US-10-161-088-1
                    66.8%; Score 349.4; DB 15; Length 651;
 Query Match
  Best Local Similarity 82.8%; Pred. No. 3e-104;
 Matches 425; Conservative
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           139 GGACCAGAGACCCTTTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGACCG 198
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           199 AGGGGCTTTTACTTCAACAAGCCCACAGGCTATGGCTCCAGCATTCGGAGGGCACCTCAG 258
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           259 ACAGGCATTGTGGATGAGTGTTGCTTCCGGAGCTGTGATCTGAGGAGACTGGAGATGTAC 318
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        Qу
           319 TGTGCCCCACTGAAGCCTACAAAAGCAGCCCGCTCTATCCGTGCCCAGCGCCACACTGAC 378
Dh
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           499 CAGAATGTAGGAGGAGCCTCCCACGGAGCAGAAAATGCCACATCACCGCAGGATCCTTTG 558
Db
       421 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCA-----AAAAATAAGTTTGATC 474
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                         559 CTGCTTGAGCAACCTGCAAAACATCGAAACACCTACCAAATAACAATAATAAGTCCAATA 618
Db
       475 ACATTTCAAAGAT-GGCATTTCCCCCAATGAAA 506
Qу
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        619 ACATTACAAAGATGGGCATTTCCCCCCAATGAAA 651
RESULT 6
US-09-919-497-24
; Sequence 24, Application US/09919497
; Patent No. US20020106662A1
; GENERAL INFORMATION:
  APPLICANT: Mutter, George L.
  TITLE OF INVENTION: PROGNOSTIC CLASSIFICATION OF ENDOMETRIAL CANCER
  FILE REFERENCE: B0801/7225
  CURRENT APPLICATION NUMBER: US/09/919,497
  CURRENT FILING DATE: 2001-07-31
  PRIOR APPLICATION NUMBER: US 60/221,735
  PRIOR FILING DATE: 2000-07-31
  NUMBER OF SEQ ID NOS: 100
  SOFTWARE: PatentIn version 3.0
 SEQ ID NO 24
   LENGTH: 7260
   TYPE: DNA
   ORGANISM: Homo sapiens
US-09-919-497-24
                    63.9%; Score 334.4; DB 10; Length 7260;
 Query Match
 Best Local Similarity
                    84.6%; Pred. No. 9e-99;
 Matches 445; Conservative
                         0; Mismatches
                                       26; Indels
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Ov
           311 GGACCGGAGACGCTCTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGGAGAC 370
Db
         61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
Qу
           Db
        371 AGGGGCTTTTATTTCAACAAGCCCACAGGGTATGGCTCCAGCAGTCGGAGGGCGCCTCAG 430
        121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
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           431 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTAAGGAGGCTGGAGATGTAT 490
Db
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Db
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           Db
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           Db
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        479 TTCAAAGAT-GGCATTTCCCCCCAATGAAATACACAAGTAAACATTC 523
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        739 TTAAAAGATGGGCGTTTCCCCCCAATGAAATACACAAGTAAACATTC 784
RESULT 7
US-09-880-107-3739
; Sequence 3739, Application US/09880107
; Patent No. US20020142981A1
: GENERAL INFORMATION:
  APPLICANT: Horne, Darci T.
  APPLICANT: Vockley, Joseph G.
  APPLICANT: Scherf, Uwe
  APPLICANT: Gene Logic, Inc.
  TITLE OF INVENTION: Gene Expression Profiles in Liver Cancer
  FILE REFERENCE: 44921-5028-WO
  CURRENT APPLICATION NUMBER: US/09/880,107
  CURRENT FILING DATE: 2001-06-14
  PRIOR APPLICATION NUMBER: US 60/211,379
  PRIOR FILING DATE: 2000-06-14
  PRIOR APPLICATION NUMBER: US 60/237,054
  PRIOR FILING DATE: 2000-10-02
  NUMBER OF SEQ ID NOS: 3950
  SOFTWARE: PatentIn Ver. 2.1
 SEQ ID NO 3739
   LENGTH: 7260
   TYPE: DNA
   ORGANISM: Homo sapiens
   FEATURE:
   OTHER INFORMATION: Genbank Accession No. US20020142981A1 X57025
US-09-880-107-3739
                     63.9%; Score 334.4; DB 10; Length 7260;
 Query Match
 Best Local Similarity 84.6%; Pred. No. 9e-99;
 Matches 445; Conservative 0; Mismatches 26; Indels
                                                    55; Gaps
                                                               4;
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1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 60
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           311 GGACCGGAGACGCTCTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 370
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Qу
           371 AGGGGCTTTTATTTCAACAAGCCCACAGGGTATGGCTCCAGCAGTCGGAGGGCGCCTCAG 430
Db
       121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Oy
           431 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTAAGGAGGCTGGAGATGTAT 490
Db
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Db
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           Db
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       301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
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                566 -----AAGGAAGTACATTTGAAGAACGCAAGTAGAGGGAGTGCAGGAAACAAGAACTA 618
Db
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           619 CAGGATGTAGGAAGACCCTCCTGAGGAGTGAAGAGTGACATGCCACCGCAGGATCCTTTG 678
       421 CTCTGCAC-AGTTACCTG-TAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACAT 478
QУ
           679 CTCTGCACGAGTTACCTGTTAAACTTTGGAACACCTACCAAAAAATAAGTTTGATAACAT 738
Db
       479 TTCAAAGAT-GGCATTTCCCCCCAATGAAATACACAAGTAAACATTC 523
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           739 TTAAAAGATGGGCGTTTCCCCCCAATGAAATACACAAGTAAACATTC 784
Db
RESULT 8
US-09-873-319-707
; Sequence 707, Application US/09873319A
 Publication No. US20030134324A1
; GENERAL INFORMATION:
  APPLICANT: Munger, William E.
  APPLICANT: Kulkarni, Prakash
  APPLICANT: Getzenberg, Robert H.
  APPLICANT: Waga, Iwao
  APPLICANT: Yamamoto, Jun
  TITLE OF INVENTION: Identifying Drugs for and Diagnosis of Benign Prostatic
  TITLE OF INVENTION: Hyperplasia Using Gene Expression Profiles
  FILE REFERENCE: 44921-5029-US
  CURRENT APPLICATION NUMBER: US/09/873,319A
  CURRENT FILING DATE: 2001-06-05
  EARLIER APPLICATION NUMBER: US 60/223,323
  EARLIER FILING DATE: 2000-08-07
  NUMBER OF SEQ ID NOS: 755
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SOFTWARE: PatentIn Ver. 2.1

: SEO ID NO 707

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TYPE: DNA
  ORGANISM: Homo sapiens
   FEATURE:
   OTHER INFORMATION: Genbank Accession No. US20030134324A1 X57025
US-09-873-319-707
 Query Match
                   63.9%; Score 334.4; DB 13; Length 7260;
 Best Local Similarity
                  84.6%; Pred. No. 9e-99;
 Matches 445; Conservative
                      0; Mismatches 26; Indels
                                              55; Gaps
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Qy
          311 GGACCGGAGACGCTCTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGGAGAC 370
Db
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          371 AGGGGCTTTTATTTCAACAAGCCCACAGGGTATGGCTCCAGCAGTCGGAGGGCGCCTCAG 430
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Db
       Qу
          491 TGCGCACCCTCAAGCCTGCCAAGTCAGCTCGCTCTGTCCGTGCCCAGCGCCACACCGAC 550
Db
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          551 ATGCCCAAGACCCAG--
Db
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Qу
               566 -----AAGGAAGTACATTTGAAGAACGCAAGTAGAGGGAGTGCAGGAAACAAGAACTA 618
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          Db
       619 CAGGATGTAGGAAGACCCTCCTGAGGAGTGAAGAGTGACATGCCACCGCAGGATCCTTTG 678
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          Db
       679 CTCTGCACGAGTTACCTGTTAAACTTTGGAACACCTACCAAAAAATAAGTTTGATAACAT 738
       479 TTCAAAGAT-GGCATTTCCCCCAATGAAATACACAAGTAAACATTC 523
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       739 TTAAAAGATGGGCGTTTCCCCCAATGAAATACACAAGTAAACATTC 784
RESULT 9
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US-09-960-706-1066

LENGTH: 7260

- ; Sequence 1066, Application US/09960706
- ; Publication No. US20030134280A1
- ; GENERAL INFORMATION:
- ; APPLICANT: Munger, William E.
- ; TITLE OF INVENTION: Identifying Drugs for and Diagnosis of Benign Prostatic Hyperplasia Using
- ; TITLE OF INVENTION: Gene Expression Profiles

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FILE REFERENCE: 44921-5029-01US
  CURRENT APPLICATION NUMBER: US/09/960,706
  CURRENT FILING DATE: 2001-09-24
  PRIOR APPLICATION NUMBER: 60/223,323
  PRIOR FILING DATE: 2000-08-07
  PRIOR APPLICATION NUMBER: 09/873,319
  PRIOR FILING DATE: 2001-06-05
  NUMBER OF SEQ ID NOS: 1124
  SOFTWARE: PatentIn Ver. 2.1
 SEO ID NO 1066
  LENGTH: 7260
   TYPE: DNA
   ORGANISM: Homo sapiens
   OTHER INFORMATION: Genbank Accession No. US20030134280A1 X57025
US-09-960-706-1066
                   63.9%; Score 334.4; DB 13; Length 7260;
 Query Match
                   84.6%; Pred. No. 9e-99;
 Best Local Similarity
 Matches 445; Conservative
                       0; Mismatches 26; Indels
         1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 60
Qу
           311 GGACCGGAGACGCTCTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGGAGAC 370
Db
        61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
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           371 AGGGGCTTTTATTTCAACAAGCCCACAGGGTATGGCTCCAGCAGTCGGAGGGCGCCTCAG 430
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       121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Qу
           431 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTAAGGAGGCTGGAGATGTAT 490
Db
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Qу
           551 ATGCCCAAGACCCAG----
Dh
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Qу
                566 -----AAGGAAGTACATTTGAAGAACGCAAGTAGAGGGAGTGCAGGAAACAAGAACTA 618
Db
       361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
QУ
           619 CAGGATGTAGGAAGACCCTCCTGAGGAGTGAAGAGTGACATGCCACCGCAGGATCCTTTG 678
Db
        421 CTCTGCAC-AGTTACCTG-TAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACAT 478
Qу
           679 CTCTGCACGAGTTACCTGTTAAACTTTGGAACACCTACCAAAAAATAAGTTTGATAACAT 738
Dh
        479 TTCAAAGAT-GGCATTTCCCCCCAATGAAATACACAAGTAAACATTC 523
Qу
           739 TTAAAAGATGGGCGTTTCCCCCCAATGAAATACACAAGTAAACATTC 784
Db
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RESULT 10
US-10-136-639-4
; Sequence 4, Application US/10136639
Publication No. US20030072761A1
GENERAL INFORMATION:
  APPLICANT: LeBowitz, Jonathan
  TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR TARGETING PROTEINS ACROSS
THE BLOOD BRAIN
  TITLE OF INVENTION: BARRIER
  FILE REFERENCE: SYM-008
  CURRENT APPLICATION NUMBER: US/10/136,639
  CURRENT FILING DATE: 2002-09-06
  PRIOR APPLICATION NUMBER: US 60/329,650
  PRIOR FILING DATE: 2001-10-16
  NUMBER OF SEQ ID NOS: 4
  SOFTWARE: PatentIn version 3.0
 SEQ ID NO 4
   LENGTH: 7260
   TYPE: DNA
   ORGANISM: Homo sapiens
US-10-136-639-4
                    63.9%; Score 334.4; DB 15; Length 7260;
 Query Match
 Best Local Similarity 84.6%; Pred. No. 9e-99;
 Matches 445; Conservative 0; Mismatches 26; Indels
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Qу
           311 GGACCGGAGACGCTCTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 370
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QУ
           371 AGGGGCTTTTATTTCAACAAGCCCACAGGGTATGGCTCCAGCAGTCGGAGGGCGCCTCAG 430
Dh
        121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Qу
           431 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTAAGGAGGCTGGAGATGTAT 490
Dh
        Qу
           491 TGCGCACCCTCAAGCCTGCCAAGTCAGCTCGCTCTGTCCGTGCCCAGCGCCACACCGAC 550
Db
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Qу
           551 ATGCCCAAGACCCAG---
Dh
        301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
Qу
                566 -----AAGGAAGTACATTTGAAGAACGCAAGTAGAGGGAGTGCAGGAAACAAGAACTA 618
Db
        361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Qу
           619 CAGGATGTAGGAAGACCCTCCTGAGGAGTGAAGAGTGACATGCCACCGCAGGATCCTTTG 678
Db
        421 CTCTGCAC-AGTTACCTG-TAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACAT 478
Qу
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Db
       679 CTCTGCACGAGTTACCTGTTAAACTTTGGAACACCTACCAAAAAATAAGTTTGATAACAT 738
       479 TTCAAAGAT-GGCATTTCCCCCAATGAAATACACAAGTAAACATTC 523
Qу
           739 TTAAAAGATGGGCGTTTCCCCCCAATGAAATACACAAGTAAACATTC 784
Db
RESULT 11
US-10-207-655-54
; Sequence 54, Application US/10207655
; Publication No. US20030118592A1
; GENERAL INFORMATION:
  APPLICANT: Ledbetter, Jeffrey A.
  APPLICANT: Hayden-Ledbetter, Martha S.
  TITLE OF INVENTION: BINDING DOMAIN-IMMUNOGLOBULIN FUSION PROTEINS
  FILE REFERENCE: 390069.401C1
  CURRENT APPLICATION NUMBER: US/10/207,655
  CURRENT FILING DATE: 2002-07-25
  NUMBER OF SEQ ID NOS: 426
  SOFTWARE: PatentIn version 3.0
 SEQ ID NO 54
   LENGTH: 725
   TYPE: DNA
   ORGANISM: Homo sapiens
US-10-207-655-54
                   63.6%; Score 332.8; DB 15; Length 725;
 Query Match
 Best Local Similarity . 84.4%; Pred. No. 9.2e-99;
                         0; Mismatches 27; Indels
 Matches 444; Conservative
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Qу
           156 GGACCGGAGACGCTCTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTAGAGAC 215
Db
         61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
QУ
           216 AGGGGCTTTTATTTCAACAAGCCCACAGGGTATGGCTCCAGCAGTCGGAGGGCGCCTCAG 275
Dh
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Qу
           276 ACAGGTATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTAAGGAGGCTGGAGATGTAT 335
Db
        Qу
           336 TGCGCACCCCTCAAGCCTGCCAAGTCAGCTCGCTCTGTCCGTGCCCAGCGCCACACCGAC 395
Db
        241 ATGCCCAAGACTCAGAAGTATCAGCCTCCATCTACCAACAAGAAAATGAAGTCTCAGAGG 300
Оy
           396 ATGCCCAAGACCCAG----
Db
        301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
QУ
                 411 -----AAGGAAGTACATTTGAAGAACGCAAGTAGAGGGAGTGCAGGAAACAAGAACTA 463
Dh
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Qу
           464 CAGGATGTAGGAAGACCCTCCTGAGGAGTGAAGAGTGACATGCCACCGCAGGATCCTTTG 523
Dh
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421 CTCTGCAC-AGTTACCTG-TAAACATTGGAATACCGGCCAAAAAATAAGTTTGATCACAT 478
Qу
           524 CTCTGCACGAGTTACCTGTTAAACTTTGGAACACCTACCAAAAAATAAGTTTGATAACAT 583
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        479 TTCAAAGAT-GGCATTTCCCCCCAATGAAATACACAAGTAAACATTC 523
Qу
           Db
        584 TTAAAAGATGGGCGTTTCCCCCAATGAAATACACAAGTAAACATTC 629
RESULT 12
US-10-251-661-7
; Sequence 7, Application US/10251661
; Publication No. US20030166555A1
; GENERAL INFORMATION:
  APPLICANT: Alberini, Cristina M.
  APPLICANT: Bear, Mark F.
  TITLE OF INVENTION: Methods and Compositions for Regulating
  TITLE OF INVENTION: Memory Consolidation
  FILE REFERENCE: 3499.1001-003
  CURRENT APPLICATION NUMBER: US/10/251,661
  CURRENT FILING DATE: 2002-09-20
  PRIOR APPLICATION NUMBER: 60/193,614
  PRIOR FILING DATE: 2000-03-31
  PRIOR APPLICATION NUMBER: PCT/US01/10661
  PRIOR FILING DATE: 2001-04-02
  NUMBER OF SEQ ID NOS: 12
  SOFTWARE: FastSEQ for Windows Version 4.0
 SEQ ID NO 7
   LENGTH: 612
   TYPE: DNA
   ORGANISM: Homo sapiens
   FEATURE:
   NAME/KEY: CDS
   LOCATION: (151)...(564)
US-10-251-661-7
 Ouery Match
                     52.3%; Score 273.6; DB 13; Length 612;
 Best Local Similarity 83.0%; Pred. No. 2.6e-79;
 Matches 347; Conservative
                          0; Mismatches 19; Indels
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                                                              1;
         1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 60
Qу
           247 GGACCGGAGACGCTCTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGGAGAC 306
Db
         61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGCCACCTCAG 120
Qу
           307 AGGGGCTTTTATTTCAACAAGCCCACAGGGTATGGCTCCAGCAGTCGGAGGGCGCCTCAG 366
Db
        121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Qу
           367 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTAAGGAGGCTGGAGATGTAT 426
Db
        Qу
           427 TGCGCACCCCTCAAGCCTGCCAAGTCAGCTCGCTCTGTCCGTGCCCAGCGCCACACCGAC 486
Db
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Qу
                502 -----AAGGAAGTACATTTGAAGAACGCAAGTAGAGGGAGTGCAGGAAACAAGAACTA 554
Db
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           555 CAGGATGTAGGAAGACCCTCCTGAGGAGTGAAGAGTGACATGCCACCGCAGGATCCTT 612
Db
RESULT 13
US-09-852-261-11
; Sequence 11, Application US/09852261
; Patent No. US20020083477A1
; GENERAL INFORMATION:
  APPLICANT: GOLDSPINK, GEOFFREY
  APPLICANT: TERENGHI, GIORGIO
  TITLE OF INVENTION: REPAIR OF NERVE DAMAGE
  FILE REFERENCE: 117-351
  CURRENT APPLICATION NUMBER: US/09/852,261
  CURRENT FILING DATE: 2001-05-10
  PRIOR APPLICATION NUMBER: GB 0011278.9
  PRIOR FILING DATE: 2000-05-10
  NUMBER OF SEQ ID NOS: 14
  SOFTWARE: PatentIn Ver. 2.1
 SEQ ID NO 11
   LENGTH: 487
   TYPE: DNA
   ORGANISM: Rattus sp.
US-09-852-261-11
                     50.1%; Score 262; DB 9; Length 487;
Query Match
 Best Local Similarity 74.7%; Pred. No. 1.5e-75;
 Matches 396; Conservative 0; Mismatches 75; Indels
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         1 GGACCGGAGACGCTCTGCGGTGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGGAGAC 60
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Db
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Qу
           61 AGGGGCTTTTACTTCAACAAGCCCACAGTCTATGGCTCCAGCATTCGGAGGGCACCACAG 120
Db
        121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Qу
           121 ACGGCATTGTGGATGAGTGTTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
Db
        QУ
               181 TGTGTCCGCTGCAAGCCTACAAAGTCAGCTCGTTCCATCCGGGCCCAGCGCCACACTGAC 240
Db
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Qу
           241 ATGCCCAAGACTCAG-
Dh
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301 AGAAGGAAAGGAAGTACATTTGAAGAACACAAGTAGAGGGAGTGCAGGAAACAAGAACTA 360
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                -----AAGGAAGTACACTTGAAGAACACAAGTAGAGGAAGTGCAGGAAACAAGACCTA 308
Db
       361 CAGGATGTAGGAAGACCCTTCTGAGGAGTGAAGAAGGACAGGCCACCGCAGGACCCTTTG 420
Qу
                                 309 CAGAATGTAGGAGGAGCCTCCCGAGGAACAGAAAATGCCACGTCACCGCAAGATCCTTTG 368
Db
       421 CTCTGCACAGTTACCTGTAAACATTGGAATACCGGCCA-----AAAAATAAGTTTGATC 474
QУ
                                            369 CTGCTTGAGCAACCTGCAAAACATCGGAACACCTGCCAAATATCAATAATGAGTTCAATA 428
Dh
       475 ACATTCAAAGAT-GGCATTTCCCCCAATGAAATACACAAGTAAACATTC 523
Qу
           429 TCATTTCAGAGATGGGCATTTCCCTCAATGAAATACACAAGTAAACATTC 478
Db
RESULT 14
US-09-852-261-9
; Sequence 9, Application US/09852261
; Patent No. US20020083477A1
 GENERAL INFORMATION:
  APPLICANT: GOLDSPINK, GEOFFREY
  APPLICANT: TERENGHI, GIORGIO
  TITLE OF INVENTION: REPAIR OF NERVE DAMAGE
  FILE REFERENCE: 117-351
  CURRENT APPLICATION NUMBER: US/09/852,261
  CURRENT FILING DATE: 2001-05-10
  PRIOR APPLICATION NUMBER: GB 0011278.9
  PRIOR FILING DATE: 2000-05-10
  NUMBER OF SEQ ID NOS: 14
  SOFTWARE: PatentIn Ver. 2.1
 SEQ ID NO 9
   LENGTH: 318
   TYPE: DNA
   ORGANISM: Homo sapiens
US-09-852-261-9
                    45.4%; Score 237.6; DB 9;
 Query Match
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 Best Local Similarity 94.6%; Pred. No. 1.3e-67;
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                                          Indels
                                                  0; Gaps
 Matches 246; Conservative
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           1 GGACCGGAGACGCTCTGCGGGGCTGAGCTGGTGGATGCTCTTCAGTTCGTGTGTGGAGAC 60
Db
        61 AGGGGCTTTTATTTCAACAAGCCCACAGGATACGGCTCCAGCAGTCGGAGGGCACCTCAG 120
Qу
           61 AGGGGCTTTTATTTCAACAAGCCCACAGGGTATGGCTCCAGCAGTCGGAGGGCGCCTCAG 120
Db
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Qу
           121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTAAGGAGGCTGGAGATGTAT 180
Db
        Qу
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181 TGCGCACCCTCAAGCCTGCCAAGTCAGCTCGCTCTGTCCGTGCCCAGCGCCCACCGAC 240
Db
        241 ATGCCCAAGACTCAGAAGTA 260
Qу
           Db
        241 ATGCCCAAGACCCAGAAGGA 260
RESULT 15
US-10-238-114-1
; Sequence 1, Application US/10238114
; Publication No. US20030100073A1
; GENERAL INFORMATION:
  APPLICANT: Merial
  APPLICANT: ANDREONI , Christine Michele
  TITLE OF INVENTION: IGF-1 AS FELINE VACCINE ADJUVANT, IN PARTICULAR AGAINST
FELINE RETROVIRUS
  FILE REFERENCE: 454313-3165.1
  CURRENT APPLICATION NUMBER: US/10/238,114
  CURRENT FILING DATE: 2002-09-10
  PRIOR APPLICATION NUMBER: FR 01 11736
  PRIOR FILING DATE: 2001-09-11
  PRIOR APPLICATION NUMBER: US 60/318,666
  PRIOR FILING DATE: 2001-09-12
  NUMBER OF SEQ ID NOS: 20
  SOFTWARE: PatentIn version 3.1
 SEQ ID NO 1
   LENGTH: 462
   TYPE: DNA
   ORGANISM: Felis catus
US-10-238-114-1
                     43.6%; Score 228; DB 15; Length 462;
 Query Match
 Best Local Similarity 92.3%; Pred. No. 2.3e-64;
 Matches 240; Conservative 0; Mismatches
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                                       20;
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Qу
           145 GGACCAGAGACGCTCTGTGGGGCTGAGTTGGTGGACGCTCTTCAGTTCGTGTGGAGAC 204
Db
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Qу
           205 AGGGGTTTTTATTTCAACAAGCCCACGGGGTATGGCTCCAGCAGTCGGAGGGCACCTCAG 264
Db
        121 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGAGGCTGGAGATGTAC 180
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            265 ACAGGCATCGTGGATGAGTGCTGCTTCCGGAGCTGTGATCTGAGGCGGCTAGAGATGTAC 324
Db
        Qу
            325 TGTGCACCCCTCAAGCCTGCCAAGTCTGCCCGCTCAGTCCGTGCTCAGCGCCACACTGAC 384
Db
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Qу
            385 ATGCCCAAGGCTCAGAAGGA 404
Db
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Search completed: December 13, 2003, 11:56:48

Job time : 235.512 secs